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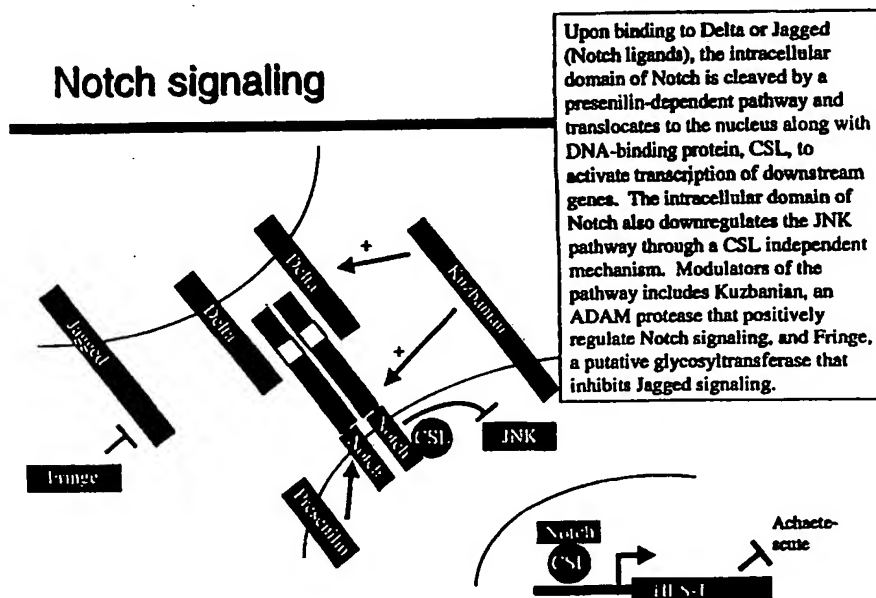
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(54) Title: NOTCH RECEPTOR LIGANDS AND USES THEREOF



(57) Abstract: The invention provides polynucleotides encoding Notch receptor ligands, encoded polypeptides, and antibodies specific to the polypeptides. Also provided are methods and compositions for enhancing or inhibiting angiogenesis as well as modulating immune responses.

NOTCH RECEPTOR LIGANDS AND USES THEREOF

FIELD OF THE INVENTION

The present invention relates to receptors and ligands consisting of human Notch gene products, and uses of these receptors, ligands and derivatives thereof to modulate cell-cell interaction in biological processes and conditions including angiogenesis and cancer.

BACKGROUND OF THE INVENTION

The Notch pathway is involved in cell fate determination and differentiation, and signaling through the Notch pathway receptors is an evolutionarily conserved mechanism for cell-cell interaction. Notch proteins, of which four have been identified in humans (Notch 1, Notch 2, Notch 3 and Notch 4), are a family of closely related transmembrane receptors. Notch 4 is expressed specifically in endothelial cells (Shiratoshi, Y., *Genes Cells* 2:213-224, 1997; Uyttendaele, H., *Development* 122:2251-2259, 1996) and may play an important role in angiogenesis. When Notch is activated by a ligand, its intracellular domain is proteolytically cleaved and transported to the nucleus, along with CSL (CBF-1/Su(H)/Lag-1/RBP-J_κ) transcription factor to activate transcription of downstream effectors. The resulting effector can repress the transcriptional activity of other genes encoding transcription factors for entry into terminal differentiation. The ligands that interact with the extracellular portion of Notch include Delta, Serrate, and Jagged; the ligands also are transmembrane proteins.

Adjacent cells of identical lineage can follow separate pathways of differentiation as a result of the Notch pathway. Sample pathways of differentiation include axis formation, cartilage formation, and somite formation. Through a process of lateral inhibition, one cell can suppress the neighboring cells from following the same path of differentiation. In one model, a Notch receptor is expressed on the cell surface of a "suppressed" cell, and interacts with a Notch ligand located on the cell surface of a dominating cell. After ligand interaction with a Notch receptor, the intracellular domain of the Notch receptor is cleaved and transported to the nucleus,

where it forms a complex and affects gene transcription. (Lendahl, U., *BioEssays* 20:103-107, 1998.)

The ligand itself plays an important role in determining the fate of cells in the vicinity of Notch-expressing cells. In *Drosophila*, ligands including Delta and Serrate have been identified and studied. The corresponding genes in mammals include Dll-1 (Delta-1) and Dll-3 (Delta-3), as well as Jag-1 and Jag-2. Mutations and decreased expression of Delta are related to phenotypic changes, and a translocation in the human Notch 1 locus (TAN-1) has been found in T-cell acute lymphoblastic leukemia/lymphoma (Ellisen et al., *Cell* 66:649-661, 1991).

10 Mutations in the human Jagged 1 gene are associated with Alagille syndrome, which involves abnormal development of liver, heart, skeleton, eye, and face. Alagille patients also exhibit valvular and arterial stenosis and high incidence of intracranial hemorrhage. Four separate mutations, all frameshifts, have been identified in patients with the syndrome. (Li, L. et al., *Nature Genetics* 16:243-251, 1997; Oda, T. et al., *Nature Genetics* 16:235-242, 1997.) The mutations are likely to interfere with the ability of Jagged 1 to interact with Notch, thereby affecting the differentiation of cells whose fate would otherwise be determined by interaction of Notch with functional Jagged 1. Mice rendered genetically deficient for Jagged 1 exhibit defects in vascular development (Xue et al., *Hum. Mol. Gen.* 8:723, 1999). These results are consistent with the hypothesis that Jagged 1 is also involved in vascular development and integrity.

Mutations in Notch 3 are related to a syndrome known as CADASIL, for cerebral autosomal arteriopathy with subcortical infarction and leukoencephalopathy. (Jontel, A. et al., *Lancet* 350:1511-1515, 1997.) Missense mutations in the extracellular domain were found in 45 out of 50 CADASIL patients in one reported study. (Salloway, S. et al., *J. Geriatr. Psychiatry Neurol.* 11:71-77, 1998.) CADASIL patients exhibit recurrent ischemic stroke and severe vascular smooth muscle cell defect. Thus, mutations in the Notch gene itself can affect vascular integrity in adults.

In view of the importance of this signaling pathway and its role in human cell differentiation and disease, there is a need in the art for identification of genes

involved in the pathway, and for methods and therapeutic agents for intervening in diseases and conditions related to defects in the Notch pathway.

SUMMARY OF THE INVENTION

The present invention relates to nucleotide sequences of human Notch
5 ligand genes, and amino acid sequences of the encoded proteins, as well as derivatives and fragments thereof wherein the derivatives and fragments exhibit biological activity such as binding to Notch receptors.

The invention also relates to methods of modulating angiogenesis by using the Notch ligands of the invention, and derivatives and fragments thereof.

10 The invention further relates to modulation of endothelial cell proliferation using polynucleotides encoding all or part of the Notch ligands of the invention, such as antisense oligonucleotides that can target nucleic acid encoding the Notch ligand.

The invention still further relates to methods of modulating the
15 development and maturation of T-cells and other cells of the immune system, thereby regulating cell-mediated immunity and antibody responses to alleviate conditions such as rheumatoid arthritis.

Notch ligands of the invention include those encoded by the 3md3 gene and the 2hd1 gene.

20 The invention relates to an isolated nucleic acid molecule comprising a polynucleotide selected from the group consisting of:

- (a) a polynucleotide encoding amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) a polynucleotide encoding amino acids from about 2 to about 583
25 of SEQ ID NO:2;
- (c) a polynucleotide encoding amino acids from about 1 to about 81 of SEQ ID NO:4;
- (d) a polynucleotide encoding amino acids from about 2 to about 81 of SEQ ID NO:4;

- (e) the polynucleotide complement of the polynucleotide of (a)-(d);
- and
- (f) a polynucleotide at least 90% identical to the polynucleotide of (a)-(e).

5 The invention also relates to an isolated nucleic acid molecule consisting of a nucleic acid comprising 50-1752 contiguous nucleotides from the coding region of SEQ ID NO:1.

 The invention further relates to an isolated nucleic acid molecule comprising a polynucleotide encoding a polypeptide wherein, except for at least one
10 conservative amino acid substitution, said polypeptide has an amino acid sequence selected from the group consisting of:

- (a) amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) amino acids from about 2 to about 583 of SEQ ID NO:2;
- (c) amino acids from about 1 to about 81 of SEQ ID NO:4; and
- 15 (d) amino acids from about 2 to about 81 of SEQ ID NO:4.

 The invention still further relates to an isolated polypeptide comprising amino acids at least 95% identical to amino acids selected from the group consisting of:

- (a) amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) amino acids from about 2 to about 583 of SEQ ID NO:2;
- 20 (c) amino acids from about 1 to about 81 of SEQ ID NO:4; and
- (d) amino acids from about 2 to about 81 of SEQ ID NO:4.

 The invention also relates to a complex comprising a protein comprising an amino acid sequence selected from the group consisting of SEQ ID NO:2 and SEQ ID NO:4.

25 The invention further relates to a complex comprising a fragment of the amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4, and a Dishevelled protein wherein said fragment is capable of forming a complex with said Dishevelled protein.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is an alignment of the amino acid sequences of novel human Notch ligand, 3md3, of the present invention (SEQ ID NO:1), the mouse ortholog Delta-3 (3721842_md3), a human Delta polypeptide (w80813_hd3), and human sequence hdelta1p.

Figure 2 is a protein phylogenic analysis based on the sequences of Figure 1, and shows that 3md3 is more closely related to murine Delta 3 than is the human sequence W80813_hd3 or human sequence hdelta1p.

Figure 3 is an alignment of the nucleotide sequences of 3md3 (SEQ ID NO:2), mouse dll3, hdelta, and W80813.

Figure 4 is a polynucleotide phylogenic analysis based on the sequences of Figure 3.

Figure 5 provides the polynucleotide (SEQ ID NO:1) and amino acid (SEQ ID NO:2) sequences of 3md3.

Figure 6 provides the polynucleotide (SEQ ID NO:3) and amino acid (SEQ ID NO:4) sequences for 2hd1.

Figure 7 is a diagram representing the interaction between the Notch receptor and the Notch ligands. Upon binding to Delta or Jagged (Notch ligands), the intracellular domain of Notch is cleaved by a presenilin-dependent pathway and translocates to the nucleus along with DNA-binding protein, CSL, to activate transcription of downstream genes. The intracellular domain of Notch also downregulates the Jun-Kinase (JNK) pathway through a CSL independent mechanism (Zecchini, V. et al., *Biol.* 9:460-499, 1999). Modulators of the pathway includes Kuzbanian, a disintegrin and metalloproteinase (ADAM) that positively regulate Notch signaling, and Fringe, a putative glycosyltransferase that inhibits Jagged signaling.

DETAILED DESCRIPTION OF THE INVENTION

Introduction

The present invention relates to nucleotide sequences of Notch ligands, the proteins encoded by the nucleotide sequences, and uses of the polynucleotides, proteins, fragments thereof, and antibodies specific for the proteins and polypeptides.

There is increasing evidence for a role of the Notch pathway in human disease. All of the components of the pathway have yet to be identified, but among those identified to date, mutations that affect their interaction with each other can lead to a variety of syndromes and pathological conditions.

A role for Notch-Jagged interaction in angiogenesis has been reported, based on studies with a model for angiogenesis, which measures bovine microvascular endothelial cell invasion into a collagen gel, through formation of a network of capillary-like tubes. (Zimrin, A. et al., *Jour. Biol. Chem.* 271:32499-32502, 1996.) Growth factors such as FGF and VEGF can be added to determine the effect of Notch and Notch ligands on growth factor-induced endothelial cell behavior and differentiation.

The Notch pathway is also implicated in the development and maturation of T cells, as described in Radtke, F. et al., *Immunity* 10:547-558, 1999. The Notch ligands of the invention are therefore useful candidates for modulating the immune system, including determining the fate of T cells, thereby affecting antibody production and/or cell-mediated immunity.

As discussed in detail below, the Notch ligands of the present invention can be used to further elucidate the role of the Notch pathway in human development and disease. In particular, the ligand gene referred to as 3md3 (SEQ ID NO:1) displays a pattern of tissue expression that overlaps with the endothelial cell-specific gene Notch 4. This has important implications for the use of 3md3 in modulating angiogenesis, such as preventing cancer-related angiogenesis to stop tumor growth and for therapeutic angiogenesis, e.g., to induce blood vessel formation in ischemia. A mouse mammary tumor virus (MMTV) oncogene, *int-3*, encodes the intracellular signaling domain of

Notch 4. Thus, the MMTV may target angiogenesis as a component of its tumor promoting activity. (Zimrin, A. et al., *J. Biol. Chem.* 271:32499-32502, 1996.)

3md3 is therefore a candidate for regulating Notch 4-mediated angiogenesis. Without being bound by a particular mechanism, applicants believe that by modulating the interaction between Notch ligand 3md3 and its receptors, the growth and differentiation of microvascular endothelial cells can be regulated. The role of the 3md3 gene product in angiogenesis can be determined using a model of angiogenesis, for example as described in more detail in the Examples.

Identification of Novel Notch Receptor Ligands

Two new ligands of the Notch receptor have been identified, and are referred to as 3md3 and 2hd1.

1. 3md3. This gene is likely the human ortholog of murine Delta 3 (Dll3, GenBank protein ID 3721842). It is expressed in the heart, kidney, skeletal muscle and kidney as a 4.9 kb message. The nucleotide sequence for the full-length coding region (1752 basepairs) is disclosed in SEQ ID NO:1, and the encoded amino acid sequence (583 amino acids) is disclosed in SEQ ID NO:2.

The mouse Dll3 gene is predominantly expressed in the neuroectoderm and paraxial mesoderm during embryogenesis (Dunwoodie, S.L. et al., *Development* 124:3065-3076, 1997). The cDNA consists of 2243 basepairs, encoding a protein of 585 amino acids. Figure 1 compares the polynucleotide sequences of 3md3 with murine Dll3. Figure 1 also compares the sequence of a human Delta polynucleotide sequence disclosed in WO 98 45434-A1 (W80813). An examination of the alignments in Figure 1 indicates that the novel sequence of the invention, 3md3, is more closely related to the mouse Delta 3 sequence than is the human Delta sequence. Therefore, it is believed that 3md3 is the human ortholog of the mouse Dll3 gene. The tissue expression of 3md3 overlaps with that of the endothelial-specific gene Notch 4, suggesting that it may be a functional ligand for Notch 4.

As described in more detail herein, the present invention provides new methods and materials for modulating angiogenesis and immune responses using products of 3md3.

2. 2hd1. This gene contains a region that is identical to the DSL (Delta/Serrate/Lag) domain in human Delta 1, followed by a divergent sequence of 24 amino acids and a stop codon. The DSL domain is a region of homology common to the known Notch ligand and is involved in receptor binding (Fitzgerald and Greenwald, *Development* 121:4275-4282, 1995). SEQ ID NO:3 represents the coding region of 2hd1, and SEQ ID NO:4 represents the translation of the coding region.

10 Polynucleotides

The invention relates to the specific polynucleotide sequences disclosed in SEQ ID NO:1 and 3, and to additional embodiments described herein. The polynucleotides of the invention also include polynucleotides having sequence similarity or sequence identity to SEQ ID NO:1 or 3. Nucleic acids having sequence similarity are detected by hybridization under low stringency conditions, for example, at 15 50°C and 10XSSC (0.9 M saline/0.09 M sodium citrate) and remain bound when subjected to washing at 55°C in 1XSSC. Sequence identity can be determined by hybridization under stringent conditions, for example, at 50°C or higher and 0.1XSSC (9 mM saline/0.9 mM sodium citrate). Hybridization methods and conditions are well known in the art, see, *e.g.*, U.S. Patent No. 5,707,829. Nucleic acids that are 20 substantially identical to the provided polynucleotide sequences, *e.g.*, allelic variants, genetically altered versions of the gene, *etc.*, bind to one of the provided polynucleotide sequences (SEQ ID NOs:1 and 3) under stringent hybridization conditions. By using probes, particularly labeled probes of DNA sequences, one of skill can isolate 25 homologous or related genes. The source of homologous genes can be any species, *e.g.*, primate species, particularly human; rodents, such as rats and mice; canines, felines, bovines, ovines, equines, yeast, nematodes, *etc.*

Preferably, hybridization is performed using at least 15 contiguous nucleotides (nt) of at least one of SEQ ID NOs:1 and 3. That is, when at least 15

contiguous nt of one of the disclosed SEQ ID NOs. is used as a probe, the probe will preferentially hybridize with a nucleic acid comprising the complementary sequence, allowing the identification and retrieval of the nucleic acids that uniquely hybridize to the selected probe. Probes of more than 15 nt can be used, *e.g.*, probes of from about 18
5 nt to about 100 nt, but 15 nt represents sufficient sequence for unique identification.

The polynucleotides of the invention also include naturally occurring variants of the nucleotide sequences (*e.g.*, degenerate variants, allelic variants). Variants of the polynucleotides of the invention are identified by hybridization of putative variants with nucleotide sequences disclosed herein, preferably by
10 hybridization under stringent conditions. For example, by using appropriate wash conditions, variants of the polynucleotides of the invention can be identified where the allelic variant exhibits at most about 25-30% base pair (bp) mismatches relative to the selected polynucleotide probe. In general, allelic variants contain 15-25% bp mismatches, and can contain as little as even 5-15%, or 2-5%, or 1-2% bp mismatches,
15 as well as a single bp mismatch.

The invention also encompasses homologs corresponding to the polynucleotides of SEQ ID NOs:1 and 3, where the source of homologous genes can be any mammalian species, *e.g.*, primate species, particularly human; rodents, such as rats; canines, felines, bovines, ovines, equines, yeast, nematodes, *etc.* Between mammalian
20 species, *e.g.*, human and mouse, homologs generally have substantial sequence similarity, *e.g.*, at least 75% sequence identity, usually at least 90%, more usually at least 95% between nucleotide sequences. Sequence similarity is calculated based on a reference sequence, which may be a subset of a larger sequence, such as a conserved motif, coding region, flanking region, *etc.* A reference sequence will usually be at least
25 about 18 contiguous nt long, more usually at least about 30 nt long, and may extend to the complete sequence that is being compared. Algorithms for sequence analysis are known in the art, such as BLAST, described in Altschul et al., *J. Mol. Biol.* 215:403-10 (1990). For the purposes of this invention, a preferred method of calculating percent identity is the Smith-Waterman algorithm, using the following. Global DNA sequence
30 identity is greater than 65% as determined by the Smith-Waterman homology search

algorithm as implemented in MPSRCH program (Oxford Molecular) using an affine gap search with the following search parameters: gap open penalty, 12; and gap extension penalty, 1.

The subject nucleic acids can be cDNAs or genomic DNAs, as well as
5 fragments thereof, particularly fragments that encode a biologically active gene product and/or are useful in the methods disclosed herein (*e.g.*, in diagnosis, as a unique identifier of a differentially expressed gene of interest, *etc.*). The term "cDNA" as used herein is intended to include all nucleic acids that share the arrangement of sequence elements found in native mature mRNA species, where sequence elements are exons
10 and 3' and 5' non-coding regions. Normally mRNA species have contiguous exons, with the intervening introns, when present, being removed by nuclear RNA splicing, to create a continuous open reading frame encoding a polypeptide of the invention.

A genomic sequence of interest comprises the nucleic acid present between the initiation codon and the stop codon, as defined in the listed sequences,
15 including all of the introns that are normally present in a native chromosome. It can also include the 3' and 5' untranslated regions found in the mature mRNA. It can further include specific transcriptional and translational regulatory sequences, such as promoters, enhancers, *etc.*, including about 1 kb, but possibly more, of flanking genomic DNA at either the 5' or 3' end of the transcribed region. The genomic DNA
20 can be isolated as a fragment of 100 kbp or smaller; and substantially free of flanking chromosomal sequence. The genomic DNA flanking the coding region, either 3' and 5', or internal regulatory sequences as sometimes found in introns, contains sequences required for proper tissue, stage-specific, or disease-state specific expression.

The nucleic acid compositions of the subject invention can encode all or
25 a part of the subject polypeptides. Double or single stranded fragments can be obtained from the DNA sequence by chemically synthesizing oligonucleotides in accordance with conventional methods, by restriction enzyme digestion, by PCR amplification, *etc.* Isolated polynucleotides and polynucleotide fragments of the invention comprise at least about 10, about 15, about 20, about 35, about 50, about 100, about 150 to about
30 200, about 250 to about 300, or about 350 contiguous nt selected from the

polynucleotide sequences as shown in SEQ ID NOs:1 and 3. Such fragments are exemplary only, and include all intervening sizes, such as 11, 12, 13, etc.; 51, 52, 53, etc.; 151, 152, 153, etc., and so on. For the most part, fragments will be of at least 15 nt, usually at least 18 nt or 25 nt, and up to at least about 50 contiguous nt in length
5 or more. In a preferred embodiment, the polynucleotide molecules comprise a contiguous sequence of at least 12 nt selected from the group consisting of the polynucleotides shown in SEQ ID NOs:1 and 3.

Preferred fragments are those which contain an antigenic determinant, and/or which are functionally active. "Functionally active" fragments include those
10 with adhesive properties. Preferred fragments of 3md3 and/or 2hd1 include the full-length protein, an extracellular domain with a transmembrane region, and fragments of the proteins that are homologous or functionally equivalent to the adhesive fragments of *Drosophila* Delta mediating heterotypic (amino acids 1-230) or homotypic (32-320) interactions. Such fragments can be expressed independently or as fusion proteins.
15 Fusion proteins may include myc-, HA-, or His6- tags. Fusion proteins may contain the Fc domain of human IgG.

Derivatives of Delta proteins, and assays for measuring the biological activity of the derivatives, are disclosed in WO 97/06571, which is incorporated by reference. WO 97/06571 also discloses therapeutics, including antisense compositions,
20 based on or derived from polynucleotides encoding Delta proteins. Such methods and compositions are applicable to the novel ligands of the invention. Truncations of Notch ligands are disclosed in U.S. Patent No. 5,648,464, which is incorporated by reference.

Probes specific to the polynucleotides of the invention can be generated using the polynucleotide sequences disclosed in SEQ ID NOs:1 and 3. The probes are
25 preferably at least about a 12, 15, 16, 18, 20, 22, 24, or 25 nt fragment of a corresponding contiguous sequence of SEQ ID NOs:1 and 3, and can be less than 2, 1, 0.5, 0.1, or 0.05 kb in length. The probes can be synthesized chemically or can be generated from longer polynucleotides using restriction enzymes. The probes can be labeled, for example, with a radioactive, biotinylated, or fluorescent tag.

The polynucleotides of the subject invention are isolated and obtained in substantial purity, generally as other than an intact chromosome. Usually, the polynucleotides, either as DNA or RNA, will be obtained substantially free of other naturally-occurring nucleic acid sequences, generally being at least about 50%, usually
5 at least about 90% pure and are typically "recombinant," *e.g.*, flanked by one or more nucleotides with which it is not normally associated on a naturally occurring chromosome.

The polynucleotides of the invention can be provided as a linear molecule or within a circular molecule, and can be provided within autonomously
10 replicating molecules (vectors) or within molecules without replication sequences. Expression of the polynucleotides can be regulated by their own or by other regulatory sequences known in the art. The polynucleotides of the invention can be introduced into suitable host cells using a variety of techniques available in the art, such as transferrin polycation-mediated DNA transfer, transfection with naked or encapsulated
15 nucleic acids, liposome-mediated DNA transfer, intracellular transportation of DNA-coated latex beads, protoplast fusion, viral infection, electroporation, gene gun, calcium phosphate-mediated transfection, and the like.

The subject nucleic acid compositions can be used, for example, to produce polypeptides, as probes for the detection of mRNA of the invention in
20 biological samples (*e.g.*, extracts of human cells), to generate additional copies of the polynucleotides, to generate ribozymes or antisense oligonucleotides, and as single stranded DNA probes or as triple-strand forming oligonucleotides. The probes described herein can be used to, for example, determine the presence or absence of the polynucleotide sequences as shown in SEQ ID NOs:1 and 3 or variants thereof in a
25 sample. These and other uses are described in more detail below.

Polypeptide Fragments

The invention provides polypeptide fragments of the disclosed protein. Polypeptide fragments of the invention can comprise at least 8, 10, 12, 15, 18, 19, 20, 25, 50, 75, 100, 125, 130, 140, 150, 160, 170, 180, 190, 200, 250, 300, 350, 400, 450,

500, 550, 560, 570 or 580 contiguous amino acids from SEQ ID NO:2. Also included are all intermediate length fragments in this range, such as 101, 102, 103, etc.; 170, 171, 172, etc.; and 600, 601, 601, etc. The specific lengths listed herein are exemplary only and not limiting. The invention also provides fragments of at least 81 contiguous amino acids from SEQ ID NO:4.

Biologically Active Variants

Variants of the protein and polypeptides disclosed herein can also occur. Variants can be naturally or non-naturally occurring. Naturally occurring variants are found in humans or other species and comprise amino acid sequences which are substantially identical to the amino acid sequence shown in SEQ ID NO:2 or 4. Species homologs of the protein can be obtained using subgenomic polynucleotides of the invention, as described below, to make suitable probes or primers to screening cDNA expression libraries from other species, such as mice, monkeys, yeast, or bacteria, identifying cDNAs which encode homologs of the protein, and expressing the cDNAs as is known in the art.

Non-naturally occurring variants which retain substantially the same biological activities as naturally occurring protein variants, specifically the four transmembrane configuration and the interaction with other cell surface proteins, are also included here. Preferably, naturally or non-naturally occurring variants have amino acid sequences which are at least 85%, 90%, or 95% identical to the amino acid sequence shown in SEQ ID NO:2 or 4. More preferably, the molecules are at least 98% or 99% identical. Percent identity is determined using any method known in the art. A non-limiting example is the Smith-Waterman homology search algorithm using an affine gap search with a gap open penalty of 12 and a gap extension penalty of 1. The Smith-Waterman homology search algorithm is taught in Smith and Waterman, *Adv. Appl. Math.* (1981) 2:482-489.

Guidance in determining which amino acid residues can be substituted, inserted, or deleted without abolishing biological or immunological activity can be found using computer programs well known in the art, such as DNASTAR software.

Preferably, amino acid changes in the protein variants disclosed herein are conservative amino acid changes, *i.e.*, substitutions of similarly charged or uncharged amino acids. A conservative amino acid change involves substitution of one of a family of amino acids which are related in their side chains. Naturally occurring amino acids are generally divided into four families: acidic (aspartate, glutamate), basic (lysine, arginine, histidine), non-polar (alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), and uncharged polar (glycine, asparagine, glutamine, cystine, serine, threonine, tyrosine) amino acids. Phenylalanine, tryptophan, and tyrosine are sometimes classified jointly as aromatic amino acids.

It is reasonable to expect that an isolated replacement of a leucine with an isoleucine or valine, an aspartate with a glutamate, a threonine with a serine, or a similar replacement of an amino acid with a structurally related amino acid will not have a major effect on the biological properties of the resulting variant.

Variants of the 3md3 and/or 2hd1 proteins disclosed herein include glycosylated forms, aggregative conjugates with other molecules, and covalent conjugates with unrelated chemical moieties. Covalent variants can be prepared by linking functionalities to groups which are found in the amino acid chain or at the N- or C-terminal residue, as is known in the art. Variants also include allelic variants, species variants, and muteins. Truncations or deletions of regions which do not affect functional activity of the proteins are also variants.

A subset of mutants, called muteins, is a group of polypeptides in which neutral amino acids, such as serines, are substituted for cysteine residues which do not participate in disulfide bonds. These mutants may be stable over a broader temperature range than native secreted proteins. See Mark *et al.*, *U.S. Patent* 4,959,314.

Preferably, amino acid changes in the 3md3 and/or 2hd1 protein or polypeptide variants are conservative amino acid changes, *i.e.*, substitutions of similarly charged or uncharged amino acids. A conservative amino acid change involves substitution of one of a family of amino acids which are related in their side chains. Naturally occurring amino acids are generally divided into four families: acidic (aspartate, glutamate), basic (lysine, arginine, histidine), non-polar (alanine, valine,

leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), and uncharged polar (glycine, asparagine, glutamine, cystine, serine, threonine, tyrosine) amino acids. Phenylalanine, tryptophan, and tyrosine are sometimes classified jointly as aromatic amino acids.

5 It is reasonable to expect that an isolated replacement of a leucine with an isoleucine or valine, an aspartate with a glutamate, a threonine with a serine, or a similar replacement of an amino acid with a structurally related amino acid will not have a major effect on the biological properties of the resulting secreted protein or polypeptide variant. Properties and functions of 3md3 and/or 2hd1 protein or
10 polypeptide variants are of the same type as a protein comprising the amino acid sequence encoded by the nucleotide sequence shown in SEQ ID NO:2 or 4, respectively, although the properties and functions of variants can differ in degree.

3md3 and/or 2hd1 protein variants include glycosylated forms, aggregative conjugates with other molecules, and covalent conjugates with unrelated
15 chemical moieties. 3md3 and/or 2hd1 protein variants also include allelic variants, species variants, and muteins. Truncations or deletions of regions which do not affect the differential expression of the 3md3 and/or 2hd1 genes are also variants. Covalent variants can be prepared by linking functionalities to groups which are found in the amino acid chain or at the N- or C-terminal residue, as is known in the art.

20 It will be recognized in the art that some amino acid sequence of the 3md3 and/or 2hd1 proteins of the invention can be varied without significant effect on the structure or function of the protein. If such differences in sequence are contemplated, it should be remembered that there are critical areas on the protein which determine activity. In general, it is possible to replace residues that form the tertiary
25 structure, provided that residues performing a similar function are used. In other instances, the type of residue may be completely unimportant if the alteration occurs at a non-critical region of the protein. The replacement of amino acids can also change the selectivity of binding to cell surface receptors. Ostade et al., *Nature* 361:266-268 (1993) describes certain mutations resulting in selective binding of TNF-alpha to only
30 one of the two known types of TNF receptors. Thus, the polypeptides of the present

invention may include one or more amino acid substitutions, deletions or additions, either from natural mutations or human manipulation.

The invention further includes variations of the 3md3 and/or 2hd1 polypeptides which show comparable expression patterns or which include antigenic regions. Such mutants include deletions, insertions, inversions, repeats, and type substitutions. Guidance concerning which amino acid changes are likely to be phenotypically silent can be found in Bowie, J.U., et al., "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions," *Science* 247:1306-1310 (1990).

Of particular interest are substitutions of charged amino acids with another charged amino acid and with neutral or negatively charged amino acids. The latter results in proteins with reduced positive charge to improve the characteristics of the disclosed protein. The prevention of aggregation is highly desirable. Aggregation of proteins not only results in a loss of activity but can also be problematic when preparing pharmaceutical formulations, because they can be immunogenic. (Pinckard et al., *Clin. Exp. Immunol.* 2:331-340 (1967); Robbins et al., *Diabetes* 36:838-845 (1987); Cleland et al., *Crit. Rev. Therapeutic Drug Carrier Systems* 10:307-377 (1993)).

Amino acids in the polypeptides of the present invention that are essential for function can be identified by methods known in the art, such as site-directed mutagenesis or alanine-scanning mutagenesis (Cunningham and Wells, *Science* 244:1081-1085 (1989)). The latter procedure introduces single alanine mutations at every residue in the molecule. The resulting mutant molecules are then tested for biological activity such as binding to a natural or synthetic binding partner. Sites that are critical for ligand-receptor binding can also be determined by structural analysis such as crystallization, nuclear magnetic resonance or photoaffinity labeling (Smith et al., *J. Mol. Biol.* 224:899-904 (1992) and de Vos et al. *Science* 255:306-312 (1992)).

As indicated, changes are preferably of a minor nature, such as conservative amino acid substitutions that do not significantly affect the folding or activity of the protein. Of course, the number of amino acid substitutions a skilled artisan would make depends on many factors, including those described above.

Generally speaking, the number of substitutions for any given polypeptide will not be more than 50, 40, 30, 25, 20, 15, 10, 5 or 3.

Fusion Proteins

Fusion proteins comprising proteins or polypeptide fragments of 3md3 and/or 2hd1 can also be constructed. Fusion proteins are useful for generating antibodies against amino acid sequences and for use in various assay systems. For example, fusion proteins can be used to identify proteins which interact with a protein of the invention or which interfere with its biological function. Physical methods, such as protein affinity chromatography, or library-based assays for protein-protein interactions, such as the yeast two-hybrid or phage display systems, can also be used for this purpose. Such methods are well known in the art and can also be used as drug screens. Fusion proteins comprising a signal sequence and/or a transmembrane domain of 3md3 and/or 2hd1 or a fragment thereof can be used to target other protein domains to cellular locations in which the domains are not normally found, such as bound to a cellular membrane or secreted extracellularly.

A fusion protein comprises two protein segments fused together by means of a peptide bond. Amino acid sequences for use in fusion proteins of the invention can be utilize the amino acid sequence shown in SEQ ID NO:2 or 4 or can be prepared from biologically active variants of SEQ ID NO:2 or 4, such as those described above. The first protein segment can include of a full-length 3md3 and/or 2hd1.

Other first protein segments can consist of at least 8, 10, 12, 15, 18, 19, 20, 25, 50, 75, 100, 125, 130, 140, 150, 160, 170, 180, 190, 200, 250, 300, 350, 400, 450, 500, 550, 560, 570 or 580 contiguous amino acids from SEQ ID NO:2.

The second protein segment can be a full-length protein or a polypeptide fragment. Proteins commonly used in fusion protein construction include β -galactosidase, β -glucuronidase, green fluorescent protein (GFP), autofluorescent proteins, including blue fluorescent protein (BFP), glutathione-S-transferase (GST), luciferase, horseradish peroxidase (HRP), and chloramphenicol acetyltransferase

(CAT). Additionally, epitope tags can be used in fusion protein constructions, including histidine (His) tags, FLAG tags, influenza hemagglutinin (HA) tags, Myc tags, VSV-G tags, and thioredoxin (Trx) tags. Other fusion constructions can include maltose binding protein (MBP), S-tag, Lex a DNA binding domain (DBD) fusions, GAL4 DNA binding domain fusions, and herpes simplex virus (HSV) BP16 protein fusions.

These fusions can be made, for example, by covalently linking two protein segments or by standard procedures in the art of molecular biology. Recombinant DNA methods can be used to prepare fusion proteins, for example, by making a DNA construct which comprises a coding sequence of SEQ ID NO:1 or 3 in proper reading frame with a nucleotide encoding the second protein segment and expressing the DNA construct in a host cell, as is known in the art. Many kits for constructing fusion proteins are available from companies that supply research labs with tools for experiments, including, for example, Promega Corporation (Madison, WI), Stratagene (La Jolla, CA), Clontech (Mountain View, CA), Santa Cruz Biotechnology (Santa Cruz, CA), MBL International Corporation (MIC; Watertown, MA), and Quantum Biotechnologies (Montreal, Canada; 1-888-DNA-KITS).

Isolation and Production of 3md3

3md3 is expressed in human microvascular endothelial cells and can be extracted from these cells or from other human cells, such as recombinant cells comprising SEQ ID NO:1 using standard biochemical methods. These methods include, but are not limited to, size exclusion chromatography, ammonium sulfate fractionation, ion exchange chromatography, affinity chromatography, crystallization, electrofocusing, and preparative gel electrophoresis. The isolated and purified protein or polypeptide is separated from other compounds which normally associate with the protein or polypeptide in a cell, such as other proteins, carbohydrates, lipids, or subcellular organelles. A preparation of isolated and purified protein or polypeptide is at least 80% pure; preferably, the preparations are 90%, 95%, or 99% pure. Purity of the preparations can be assessed by any means known in the art. For example, the

purity of a preparation can be assessed by examining electrophoretograms of protein or polypeptide preparations at several pH values and at several polyacrylamide concentrations, as is known in the art.

Proteins, fusion proteins, or polypeptides of the invention can be produced by recombinant DNA methods. For production of recombinant proteins, fusion proteins, or polypeptides, a coding sequence of the nucleotide sequence shown in SEQ ID NO:1 or 3 can be expressed in prokaryotic or eukaryotic host cells using expression systems known in the art. These expression systems include bacterial, yeast, insect, and mammalian cells.

The resulting expressed 3md3 and/or 2hd1 proteins can then be purified from the culture medium or from extracts of the cultured cells using purification procedures known in the art. For example, for proteins fully secreted into the culture medium, cell-free medium can be diluted with sodium acetate and contacted with a cation exchange resin, followed by hydrophobic interaction chromatography. Using this method, the desired protein or polypeptide is typically greater than 95% pure. Further purification can be undertaken, using, for example, any of the techniques listed above.

It may be necessary to modify a protein produced in yeast or bacteria, for example by phosphorylation or glycosylation of the appropriate sites, in order to obtain a functional protein. Such covalent attachments can be made using known chemical or enzymatic methods.

3md3 and/or 2hd1 protein or polypeptide of the invention can also be expressed in cultured host cells in a form which will facilitate purification. For example, a protein or polypeptide can be expressed as a fusion protein comprising, for example, maltose binding protein, glutathione-S-transferase, or thioredoxin, and purified using a commercially available kit. Kits for expression and purification of such fusion proteins are available from companies such as New England BioLabs, Pharmacia, and Invitrogen. Proteins, fusion proteins, or polypeptides can also be tagged with an epitope, such as a "Flag" epitope (Kodak), and purified using an antibody which specifically binds to that epitope.

The coding sequence disclosed herein can also be used to construct transgenic animals, such as cows, goats, pigs, or sheep. Female transgenic animals can then produce proteins, polypeptides, or fusion proteins of the invention in their milk. Methods for constructing such animals are known and widely used in the art.

5 Alternatively, synthetic chemical methods, such as solid phase peptide synthesis, can be used to synthesize a secreted protein or polypeptide. General means for the production of peptides, analogs or derivatives are outlined in Chemistry and Biochemistry of Amino Acids, Peptides, and Proteins -- A Survey of Recent Developments, B. Weinstein, ed. (1983). Substitution of D-amino acids for the normal
10 L-stereoisomer can be carried out to increase the half-life of the molecule. Variants can be similarly produced.

 Isolated genes corresponding to the cDNA sequences disclosed herein are also provided. Standard molecular biology methods can be used to isolate the corresponding genes using the cDNA sequences provided herein. These methods
15 include preparation of probes or primers from the nucleotide sequence shown in SEQ ID NO:1 or 3 for use in identifying or amplifying the genes from human genomic libraries or other sources of human genomic DNA.

 Polynucleotide molecules of the invention can also be used as primers to obtain additional copies of the polynucleotides, using polynucleotide amplification
20 methods. Polynucleotide molecules can be propagated in vectors and cell lines using techniques well known in the art. Polynucleotide molecules can be on linear or circular molecules. They can be on autonomously replicating molecules or on molecules without replication sequences. They can be regulated by their own or by other regulatory sequences, as is known in the art.

25 Polynucleotide Constructs

 Polynucleotide molecules comprising the coding sequences disclosed herein can be used in a polynucleotide construct, such as a DNA or RNA construct. Polynucleotide molecules of the invention can be used, for example, in an expression construct to express all or a portion of a secreted protein, variant, fusion protein, or

single-chain antibody in a host cell. An expression construct comprises a promoter which is functional in a chosen host cell. The skilled artisan can readily select an appropriate promoter from the large number of cell type-specific promoters known and used in the art. The expression construct can also contain a transcription terminator
 5 which is functional in the host cell. The expression construct comprises a polynucleotide segment which encodes all or a portion of the desired protein. The polynucleotide segment is located downstream from the promoter. Transcription of the polynucleotide segment initiates at the promoter. The expression construct can be linear or circular and can contain sequences, if desired, for autonomous replication.

10 Host Cells

An expression construct can be introduced into a host cell. The host cell comprising the expression construct can be any suitable prokaryotic or eukaryotic cell. Expression systems in bacteria include those described in Chang *et al.*, *Nature* (1978) 275: 615; Goeddel *et al.*, *Nature* (1979) 281: 544; Goeddel *et al.*, *Nucleic Acids Res.*
 15 (1980) 8: 4057; EP 36,776; U.S. 4,551,433; deBoer *et al.*, *Proc. Natl. Acad. Sci. USA* (1983) 80: 21-25; and Siebenlist *et al.*, *Cell* (1980) 20: 269.

Expression systems in yeast include those described in Hinnen *et al.*, *Proc. Natl. Acad. Sci. USA* (1978) 75: 1929; Ito *et al.*, *J. Bacteriol.* (1983) 153: 163; Kurtz *et al.*, *Mol. Cell. Biol.* (1986) 6: 142; Kunze *et al.*, *J Basic Microbiol.* (1985) 25:
 20 141; Gleeson *et al.*, *J. Gen. Microbiol.* (1986) 132: 3459, Roggenkamp *et al.*, *Mol. Gen. Genet.* (1986) 202 :302); Das *et al.*, *J Bacteriol.* (1984) 158: 1165; De Louvencourt *et al.*, *J. Bacteriol.* (1983) 154: 737, Van den Berg *et al.*, *Bio/Technology* (1990) 8: 135; Kunze *et al.*, *J. Basic Microbiol.* (1985) 25: 141; Cregg *et al.*, *Mol. Cell. Biol.* (1985) 5: 3376; U.S. 4,837,148; U.S. 4,929,555; Beach and Nurse, *Nature* (1981) 300: 706;
 25 Davidow *et al.*, *Curr. Genet.* (1985) 1p: 380; Gaillardin *et al.*, *Curr. Genet.* (1985) 10: 49; Ballance *et al.*, *Biochem. Biophys. Res. Commun.* (1983) 112: 284-289; Tilburn *et al.*, *Gene* (1983) 26: 205-22; Yelton *et al.*, *Proc. Natl. Acad. Sci. USA* (1984) 81: 1470-1474; Kelly and Hynes, *EMBO J.* (1985) 4: 475479; EP 244,234; and WO 91/00357.

Expression of heterologous genes in insects can be accomplished as described in U.S. 4,745,051; Friesen *et al.* (1986) "The Regulation of Baculovirus Gene Expression" in: THE MOLECULAR BIOLOGY OF BACULOVIRUSES (W. Doerfler, ed.); EP 127,839; EP 155,476; Vlak *et al.*, *J. Gen. Virol.* (1988) 69: 765-776; Miller *et al.*, *Ann. Rev. Microbiol.* (1988) 42: 177; Carbonell *et al.*, *Gene* (1988) 73: 409; Maeda *et al.*, *Nature* (1985) 315: 592-594; Lebacqz-Verheyden *et al.*, *Mol. Cell Biol.* (1988) 8: 3129; Smith *et al.*, *Proc. Natl. Acad. Sci. USA* (1985) 82: 8404; Miyajima *et al.*, *Gene* (1987) 58: 273; and Martin *et al.*, *DNA* (1988) 7:99. Numerous baculoviral strains and variants and corresponding permissive insect host cells from hosts are described in
10 Luckow *et al.*, *Bio/Technology* (1988) 6: 47-55, Miller *et al.*, in GENERIC ENGINEERING (Setlow, J.K. *et al.* eds.), Vol. 8 (Plenum Publishing, 1986), pp. 277-279; and Maeda *et al.*, *Nature*, (1985) 315: 592-594.

Mammalian expression can be accomplished as described in Dijkema *et al.*, *EMBO J.* (1985) 4: 761; Gorman *et al.*, *Proc. Natl. Acad. Sci. USA* (1982b) 79:
15 6777; Boshart *et al.*, *Cell* (1985) 41: 521; and U.S. 4,399,216. Other features of mammalian expression can be facilitated as described in Ham and Wallace, *Meth Enz.* (1979) 58: 44; Barnes and Sato, *Anal. Biochem.* (1980) 102: 255; U.S. 4,767,704; U.S. 4,657,866; U.S. 4,927,762; U.S. 4,560,655; WO 90/103430, WO 87/00195, and U.S. RE 30,985.

20 Expression constructs can be introduced into host cells using any technique known in the art. These techniques include transferrin-polycation-mediated DNA transfer, transfection with naked or encapsulated nucleic acids, liposome-mediated cellular fusion, intracellular transportation of DNA-coated latex beads, protoplast fusion, viral infection, electroporation, "gene gun," and calcium phosphate-
25 mediated transfection.

Expression of an endogenous gene encoding a protein of the invention can also be manipulated by introducing by homologous recombination a DNA construct comprising a transcription unit in frame with the endogenous gene, to form a homologously recombinant cell comprising the transcription unit. The transcription
30 unit comprises a targeting sequence, a regulatory sequence, an exon, and an unpaired

splice donor site. The new transcription unit can be used to turn the endogenous gene on or off as desired. This method of affecting endogenous gene expression is taught in U.S. Patent 5,641,670.

The targeting sequence is a segment of at least 10, 12, 15, 20, or 50 contiguous nucleotides from the nucleotide sequence shown in SEQ ID NO:1 or 3. The transcription unit is located upstream to a coding sequence of the endogenous gene. The exogenous regulatory sequence directs transcription of the coding sequence of the endogenous gene.

Expression of Polypeptide Encoded by Full-Length cDNA or Full-Length Gene

The provided polynucleotides (*e.g.*, a polynucleotide having a sequence of one of SEQ ID NOs:1 and 3), the corresponding cDNA, or the full-length gene is used to express a partial or complete gene product. Constructs of polynucleotides having sequences of SEQ ID NOs:1 and 3 can be generated synthetically. Alternatively, single-step assembly of a gene and entire plasmid from large numbers of oligodeoxyribonucleotides is described by, *e.g.*, Stemmer et al., *Gene (Amsterdam)* (1995) 164(1):49-53. In this method, assembly PCR (the synthesis of long DNA sequences from large numbers of oligodeoxyribonucleotides) is derived from DNA shuffling (Stemmer, *Nature* (1994) 370:389-391).

Appropriate polynucleotide constructs are purified using standard recombinant DNA techniques as described in, for example, Sambrook et al., *Molecular Cloning: A Laboratory Manual, 2nd Ed.*, (1989) Cold Spring Harbor Press, Cold Spring Harbor, NY. The gene product encoded by a polynucleotide of the invention is expressed in any expression system, including, for example, bacterial, yeast, insect, amphibian and mammalian systems. Vectors, host cells and methods for obtaining expression in same are well known in the art. Suitable vectors and host cells are described in U.S. Patent No. 5,654,173.

Polynucleotide molecules comprising a polynucleotide sequence provided herein are generally propagated by placing the molecule in a vector. Viral and non-viral vectors are used, including plasmids. The choice of plasmid will depend on

the type of cell in which propagation is desired and the purpose of propagation. Certain vectors are useful for amplifying and making large amounts of the desired DNA sequence. Other vectors are suitable for expression in cells in culture. Still other vectors are suitable for transfer and expression in cells in a whole animal or person.

- 5 The choice of appropriate vector is well within the skill of the art. Many such vectors are available commercially. Methods for preparation of vectors comprising a desired sequence are well known in the art.

- The polynucleotides set forth in SEQ ID NOs:1 and 3 or their corresponding full-length polynucleotides are linked to regulatory sequences as
10 appropriate to obtain the desired expression properties. These can include promoters (attached either at the 5' end of the sense strand or at the 3' end of the antisense strand), enhancers, terminators, operators, repressors, and inducers. The promoters can be regulated or constitutive. In some situations it may be desirable to use conditionally active promoters, such as tissue-specific or developmental stage-specific promoters.
15 These are linked to the desired nucleotide sequence using the techniques described above for linkage to vectors. Any techniques known in the art can be used.

- When any appropriate host cells or organisms are used to replicate and/or express the polynucleotides or nucleic acids of the invention, the resulting replicated nucleic acid, RNA, expressed protein or polypeptide, is within the scope of
20 the invention as a product of the host cell or organism. The product is recovered by any appropriate means known in the art.

- Expression of a gene corresponding to SEQ ID NO:1 or 3 can be regulated in the cell to which the gene is native. For example, an endogenous gene of a cell can be regulated by an exogenous regulatory sequence as disclosed in U.S. Patent
25 No. 5,641,670.

Antibody Production

Expression products of a polynucleotide of the invention, as well as the corresponding mRNA, cDNA, or full gene, can be prepared and used for raising antibodies for experimental, diagnostic, and therapeutic purposes. For example, antibodies

can be generated using fusion proteins or peptide sequences derived from the corresponding protein. The polynucleotide or related cDNA is expressed as described above, and antibodies are prepared. These antibodies are specific to an epitope on the polypeptide encoded by the polynucleotide, and can precipitate or bind to the
5 corresponding native protein in a cell or tissue preparation or in a cell-free extract of an *in vitro* expression system.

Methods for production of monoclonal and polyclonal antibodies that specifically bind a selected antigen are well known in the art. Typically, at least 6, 8, 10, or 12 contiguous amino acids are required to form an epitope. Epitopes that involve
10 non-contiguous amino acids may require a longer polypeptide, *e.g.*, at least 15, 25, or 50 amino acids. Antibodies that specifically bind to human polypeptides encoded by the provided polynucleotides should provide a detection signal at least 5-, 10-, or 20-fold higher than a detection signal provided with other proteins when used in Western blots or other immunochemical assays. Preferably, antibodies that specifically bind poly-
15 peptides of the invention do not bind to other proteins in immunochemical assays at detectable levels and can immunoprecipitate the specific polypeptide from solution.

The invention also contemplates naturally occurring antibodies specific for a polypeptide of the invention. For example, serum antibodies to a polypeptide of the invention in a human population can be purified by methods well known in the art,
20 *e.g.*, by passing antiserum over a column to which the corresponding selected polypeptide or fusion protein is bound. The bound antibodies can then be eluted from the column, for example using a buffer with a high salt concentration.

In addition to the antibodies discussed above, the invention also contemplates genetically engineered antibodies, antibody derivatives (*e.g.*, single chain
25 antibodies, antibody fragments (*e.g.*, Fab, etc.)), according to methods well known in the art.

In certain embodiments of the present invention, humanized anti-3md3 or 2hd1 monoclonal antibodies are provided. The phrase "humanized antibody" refers to an antibody derived from a non-human antibody, typically a mouse monoclonal
30 antibody. Alternatively, a humanized antibody may be derived from a chimeric

antibody that retains or substantially retains the antigen-binding properties of the parental, non-human, antibody but which exhibits diminished immunogenicity as compared to the parental antibody when administered to humans. The phrase "chimeric antibody," as used herein, refers to an antibody containing sequence derived from two
5 different antibodies (*see, e.g.*, U.S. Patent No. 4,816,567) which typically originate from different species. Most typically, chimeric antibodies comprise human and murine antibody fragments, generally human constant and mouse variable regions.

Because humanized antibodies are far less immunogenic in humans than the parental mouse monoclonal antibodies, they can be used for the treatment of humans
10 with far less risk of anaphylaxis. Thus, these antibodies may be preferred in therapeutic applications that involve *in vivo* administration to a human such as, *e.g.*, use as radiation sensitizers for the treatment of neoplastic disease or use in methods to reduce the side effects of, *e.g.*, cancer therapy.

Humanized antibodies may be achieved by a variety of methods
15 including, for example: (1) grafting the non-human complementarity determining regions (CDRs) onto a human framework and constant region (a process referred to in the art as "humanizing"), or, alternatively, (2) transplanting the entire non-human variable domains, but "cloaking" them with a human-like surface by replacement of surface residues (a process referred to in the art as "veneering"). In the present
20 invention, humanized antibodies will include both "humanized" and "veneered" antibodies. These methods are disclosed in, *e.g.*, Jones et al., *Nature* 321:522-525 (1986); Morrison et al., *Proc. Natl. Acad. Sci., U.S.A.*, 81:6851-6855 (1984); Morrison and Oi, *Adv. Immunol.*, 44:65-92 (1988); Verhoeyer et al., *Science* 239:1534-1536 (1988); Padlan, *Molec. Immunol.* 28:489-498 (1991); Padlan, *Molec. Immunol.* 31(3):169-
25 217 (1994); and Kettleborough, C.A. et al., *Protein Eng.* 4(7):773-83 (1991) each of which is incorporated herein by reference.

The phrase "complementarity determining region" refers to amino acid sequences which together define the binding affinity and specificity of the natural Fv region of a native immunoglobulin binding site. *See, e.g.*, Chothia et al., *J. Mol. Biol.*
30 196:901-917 (1987); Kabat et al., U.S. Dept. of Health and Human Services NIH

Publication No. 91-3242 (1991). The phrase "constant region" refers to the portion of the antibody molecule that confers effector functions. In the present invention, mouse constant regions are substituted by human constant regions. The constant regions of the subject humanized antibodies are derived from human immunoglobulins. The heavy chain constant region can be selected from any of the five isotypes: alpha, delta, epsilon, gamma or mu.

One method of humanizing antibodies comprises aligning the non-human heavy and light chain sequences to human heavy and light chain sequences, selecting and replacing the non-human framework with a human framework based on such alignment, molecular modeling to predict the conformation of the humanized sequence and comparing to the conformation of the parent antibody. This process is followed by repeated back mutation of residues in the CDR region which disturb the structure of the CDRs until the predicted conformation of the humanized sequence model closely approximates the conformation of the non-human CDRs of the parent non-human antibody. Such humanized antibodies may be further derivatized to facilitate uptake and clearance, *e.g.*, via Ashwell receptors. *See, e.g.*, U.S. Patent Nos. 5,530,101 and 5,585,089 which patents are incorporated herein by reference.

Humanized antibodies to 3md3 and/or 2hd1 can also be produced using transgenic animals that are engineered to contain human immunoglobulin loci. For example, WO 98/24893 discloses transgenic animals having a human Ig locus wherein the animals do not produce functional endogenous immunoglobulins due to the inactivation of endogenous heavy and light chain loci. WO 91/10741 also discloses transgenic non-primate mammalian hosts capable of mounting an immune response to an immunogen, wherein the antibodies have primate constant and/or variable regions, and wherein the endogenous immunoglobulin-encoding loci are substituted or inactivated. WO 96/30498 discloses the use of the Cre/Lox system to modify the immunoglobulin locus in a mammal, such as to replace all or a portion of the constant or variable region to form a modified antibody molecule. WO 94/02602 discloses non-human mammalian hosts having inactivated endogenous Ig loci and functional human Ig loci. U.S. Patent No. 5,939,598 discloses methods of making transgenic mice in

which the mice lack endogenous heavy chains, and express an exogenous immunoglobulin locus comprising one or more xenogeneic constant regions.

Using a transgenic animal described above, an immune response can be produced to a selected antigenic molecule, and antibody-producing cells can be removed from the animal and used to produce hybridomas that secrete human monoclonal antibodies. Immunization protocols, adjuvants, and the like are known in the art, and are used in immunization of, for example, a transgenic mouse as described in WO 96/33735. This publication discloses monoclonal antibodies against a variety of antigenic molecules including IL-6, IL-8, TNF α , human CD4, L-selectin, gp39, and tetanus toxin. The monoclonal antibodies can be tested for the ability to inhibit or neutralize the biological activity or physiological effect of the corresponding protein. WO 96/33735 discloses that monoclonal antibodies against IL-8, derived from immune cells of transgenic mice immunized with IL-8, blocked IL-8-induced functions of neutrophils. Human monoclonal antibodies with specificity for the antigen used to immunize transgenic animals are also disclosed in WO 96/34096.

In the present invention, 3md3 and/or 2hd1 polypeptides of the invention and variants thereof are used to immunize a transgenic animal as described above. Monoclonal antibodies are made using methods known in the art, and the specificity of the antibodies is tested using corresponding isolated 3md3 and/or 2hd1 polypeptides.

20 Polynucleotides or Arrays for Diagnostics

Polynucleotide arrays are created by spotting polynucleotide probes onto a substrate (*e.g.*, glass, nitrocellulose, *etc.*) in a two-dimensional matrix or array having bound probes. The probes can be bound to the substrate by either covalent bonds or by non-specific interactions, such as hydrophobic interactions. Samples of polynucleotides can be detectably labeled (*e.g.*, using radioactive or fluorescent labels) and then hybridized to the probes. Double stranded polynucleotides, comprising the labeled sample polynucleotides bound to probe polynucleotides, can be detected once the unbound portion of the sample is washed away. Techniques for constructing arrays and methods of using these arrays are described in EP 799 897; WO 97/29212; WO

97/27317; EP 785 280; WO 97/02357; U.S. Patent No. 5,593,839; U.S. Patent No. 5,578,832; EP 728 520; U.S. Patent No. 5,599,695; EP 721 016; U.S. Patent No. 5,556,752; WO 95/22058; and U.S. Patent No. 5,631,734. Arrays can be used to detect differential expression of a polynucleotide between a test cell and control cell (e.g.,
5 cancer cells and normal cells). For example, high expression of a particular message in a cancer cell, which is not observed in a corresponding normal cell, can indicate a cancer specific gene product. Exemplary uses of arrays are further described in, for example, Pappalarado et al., *Sem. Radiation Oncol.* (1998) 8:217; and Ramsay, *Nature Biotechnol.* (1998) 16:40.

10 Differential Expression in Diagnosis

The polynucleotides of the invention can also be used to detect differences in expression levels between two cells, e.g., as a method to identify abnormal or diseased tissue in a human. In general, the expression of a gene corresponding to a specific polynucleotide is compared between a first tissue that is
15 suspected of being diseased and a second, normal tissue of the human. The tissue suspected of being abnormal or diseased can be derived from a different tissue type of the human, but preferably it is derived from the same tissue type; for example an intestinal polyp or other abnormal growth should be compared with normal intestinal tissue. The normal tissue can be the same tissue as that of the test sample, or any
20 normal tissue of the patient, especially those that express the polynucleotide-related gene of interest (e.g., brain, thymus, testis, heart, prostate, placenta, spleen, small intestine, skeletal muscle, pancreas, breast, ovary, lung, and the mucosal lining of the colon). A difference between the polynucleotide-related gene, mRNA, or protein in the two tissues which are compared, for example in molecular weight, amino acid or
25 nucleotide sequence, or relative abundance, indicates a change in the gene, or a gene which regulates it, in the tissue of the human that was suspected of being diseased. Examples of detection of differential expression and its use in diagnosis of cancer are described in U.S. Patent Nos. 5,688,641 and 5,677,125.

Because mutations and defects in the Notch pathway are implicated in developmental disorders, a genetic predisposition to Notch pathway-related disease in a human can also be detected by comparing expression levels of an mRNA or protein corresponding to a polynucleotide of the invention in a fetal tissue, with levels in
5 normal fetal tissue. Fetal tissues that are used for this purpose include, but are not limited to, amniotic fluid, chorionic villi, blood, and the blastomere of an *in vitro*-fertilized embryo. The comparable normal polynucleotide-related gene is obtained from any tissue. The mRNA or protein is obtained from a normal tissue of a human in which the polynucleotide-related gene is expressed. Differences such as alterations in the
10 nucleotide sequence or size of the same product of the fetal polynucleotide-related gene or mRNA, or alterations in the molecular weight, amino acid sequence, or relative abundance of fetal protein, can indicate a germline mutation in the polynucleotide-related gene of the fetus, which indicates a genetic predisposition to disease.

In general, diagnostic, prognostic, and other methods of the invention
15 based on differential expression involve detection of a level or amount of a gene product, particularly a differentially expressed gene product, in a test sample obtained from a patient suspected of having or being susceptible to a disease (*e.g.*, breast cancer, lung cancer, colon cancer and/or metastatic forms thereof), and comparing the detected levels to those levels found in normal cells (*e.g.*, cells substantially unaffected by
20 cancer) and/or other control cells (*e.g.*, to differentiate a cancerous cell from a cell affected by dysplasia). Furthermore, the severity of the disease can be assessed by comparing the detected levels of a differentially expressed gene product with those levels detected in samples representing the levels of differentially gene product associated with varying degrees of severity of disease. It should be noted that use of the
25 term "diagnostic" herein is not necessarily meant to exclude "prognostic" or "prognosis," but rather is used as a matter of convenience.

The term "differentially expressed gene" is generally intended to encompass a polynucleotide that can, for example, include an open reading frame encoding a gene product (*e.g.*, a polypeptide), and/or introns of such genes and adjacent
30 5' and 3' non-coding nucleotide sequences involved in the regulation of expression, up

to about 20 kb beyond the coding region, but possibly further in either direction. The gene can be introduced into an appropriate vector for extrachromosomal maintenance or for integration into a host genome. In general, a difference in expression level associated with a decrease in expression level of at least about 25%, usually at least about 50% to 75%, more usually at least about 90% or more is indicative of a differentially expressed gene of interest, *i.e.*, a gene that is underexpressed or down-regulated in the test sample relative to a control sample. Furthermore, a difference in expression level associated with an increase in expression of at least about 25%, usually at least about 50% to 75%, more usually at least about 90% and can be at least about 1 1/2-fold, usually at least about 2-fold to about 10-fold, or about 100-fold to about 1,000-fold increase relative to a control sample is indicative of a differentially expressed gene of interest, *i.e.*, an overexpressed or up-regulated gene.

“Differentially expressed polynucleotide” as used herein means a nucleic acid molecule (RNA or DNA) comprising a sequence that represents a differentially expressed gene, *e.g.*, the differentially expressed polynucleotide comprises a sequence (*e.g.*, an open reading frame encoding a gene product) that uniquely identifies a differentially expressed gene so that detection of the differentially expressed polynucleotide in a sample is correlated with the presence of a differentially expressed gene in a sample. “Differentially expressed polynucleotides” is also meant to encompass fragments of the disclosed polynucleotides, *e.g.*, fragments retaining biological activity, as well as nucleic acids homologous, substantially similar, or substantially identical (*e.g.*, having about 90% sequence identity) to the disclosed polynucleotides.

“Diagnosis” as used herein generally includes determination of a subject's susceptibility to a disease or disorder, determination as to whether a subject is presently affected by a disease or disorder, as well as the prognosis of a subject affected by a disease or disorder (*e.g.*, identification of pre-metastatic or metastatic cancerous states, stages of cancer, or responsiveness of cancer to therapy).

“Sample” or “biological sample” as used throughout here are generally meant to refer to samples of biological fluids or tissues, particularly samples obtained

from tissues, especially from cells of the type associated with the disease for which the diagnostic application is designed, and the like. "Samples" is also meant to encompass derivatives and fractions of such samples (*e.g.*, cell lysates). Where the sample is solid tissue, the cells of the tissue can be dissociated or tissue sections can be analyzed.

5 Methods of the subject invention useful in diagnosis or prognosis typically involve comparison of the abundance of a selected differentially expressed gene product in a sample of interest with that of a control to determine any relative differences in the expression of the gene product, where the difference can be measured qualitatively and/or quantitatively. Quantitation can be accomplished, for example, by
10 comparing the level of expression product detected in the sample with the amounts of product present in a standard curve. A comparison can be made visually; by using a technique such as densitometry, with or without computerized assistance; by preparing a representative library of cDNA clones of mRNA isolated from a test sample, sequencing the clones in the library to determine that number of cDNA clones
15 corresponding to the same gene product, and analyzing the number of clones corresponding to that same gene product relative to the number of clones of the same gene product in a control sample; or by using an array to detect relative levels of hybridization to a selected sequence or set of sequences, and comparing the hybridization pattern to that of a control. The differences in expression are then
20 correlated with the presence or absence of an abnormal expression pattern. Various methods for determining the nucleic acid abundance in a sample are known to those of skill in the art (see, *e.g.*, WO 97/27317). In general, diagnostic assays of the invention involve detection of a gene product of the polynucleotide sequence (*e.g.*, mRNA or polypeptide) that corresponds to a sequence of SEQ ID NOs:1 and 3. The patient from
25 whom the sample is obtained can be apparently healthy, susceptible to disease (*e.g.*, as determined by family history or exposure to certain environmental factors), or can already be identified as having a condition in which altered expression of a gene product of the invention is implicated.

 Diagnosis can be determined based on detected gene product expression
30 levels of a gene product encoded by at least one of the polynucleotides having a

sequence set forth in SEQ ID NOs:1 and 3, and can involve detection of expression of genes corresponding to all of SEQ ID NOs:1 and 3 and/or additional sequences that can serve as additional diagnostic markers and/or reference sequences. Where the diagnostic method is designed to detect the presence or susceptibility of a patient to cancer, the assay preferably involves detection of a gene product encoded by a gene corresponding to a polynucleotide that is differentially expressed in cancer.

Any of a variety of detectable labels can be used in connection with the various embodiments of the diagnostic methods of the invention. Suitable detectable labels include fluorochromes, (*e.g.*, fluorescein isothiocyanate (FITC), rhodamine, Texas Red, phycoerythrin, allophycocyanin, 6-carboxyfluorescein (6-FAM), 2',7'-dimethoxy-4',5'-dichloro-6-carboxyfluorescein, 6-carboxy-X-rhodamine (ROX), 6-carboxy-2',4',7',4,7-hexachlorofluorescein (HEX), 5-carboxyfluorescein (5-FAM) or N,N,N',N'-tetramethyl-6-carboxyrhodamine (TAMRA)), radioactive labels, (*e.g.*, ^{32}P , ^{35}S , ^3H , *etc.*), and the like. The detectable label can involve a two stage systems (*e.g.*, biotin-avidin, hapten-anti-hapten antibody, *etc.*)

Reagents specific for the polynucleotides and polypeptides of the invention, such as antibodies and nucleotide probes, can be supplied in a kit for detecting the presence of an expression product in a biological sample. The kit can also contain buffers or labeling components, as well as instructions for using the reagents to detect and quantify expression products in the biological sample. Exemplary embodiments of the diagnostic methods of the invention are described below in more detail.

Polypeptide detection in diagnosis. In one embodiment, the test sample is assayed for the level of a differentially expressed polypeptide. Diagnosis can be accomplished using any of a number of methods to determine the absence or presence or altered amounts of the differentially expressed polypeptide in the test sample. For example, detection can utilize staining of cells or histological sections with labeled antibodies, performed in accordance with conventional methods. Cells can be permeabilized to stain cytoplasmic molecules. In general, antibodies that specifically bind a differentially expressed polypeptide of the invention are added to a sample, and

incubated for a period of time sufficient to allow binding to the epitope, usually at least about 10 minutes. The antibody can be detectably labeled for direct detection (*e.g.*, using radioisotopes, enzymes, fluorescers, chemilumescers, and the like), or can be used in conjunction with a second stage antibody or reagent to detect binding (*e.g.*,
5 biotin with horseradish peroxidase-conjugated avidin, a secondary antibody conjugated to a fluorescent compound, *e.g.*, fluorescein, rhodamine, Texas red, *etc.*). The absence or presence of antibody binding can be determined by various methods, including flow cytometry of dissociated cells, microscopy, radiography, scintillation counting, *etc.* Any suitable alternative methods of qualitative or quantitative detection of levels or
10 amounts of differentially expressed polypeptide can be used, for example ELISA, western blot, immunoprecipitation, radioimmunoassay, *etc.*

mRNA detection. The diagnostic methods of the invention can also or alternatively involve detection of mRNA encoded by the 3md3 and/or 2hdl gene. Any suitable qualitative or quantitative methods known in the art for detecting specific
15 mRNAs can be used. mRNA can be detected by, for example, *in situ* hybridization in tissue sections, by reverse transcriptase-PCR, or in Northern blots containing poly A+ mRNA. One of skill in the art can readily use these methods to determine differences in the size or amount of mRNA transcripts between two samples. mRNA expression levels in a sample can also be determined by generation of a library of expressed
20 sequence tags (ESTs) from the sample, where the EST library is representative of sequences present in the sample (Adams, et al., (1991) *Science* 252:1651). Enumeration of the relative representation of ESTs within the library can be used to approximate the relative representation of the gene transcript within the starting sample. The results of EST analysis of a test sample can then be compared to EST analysis of a reference
25 sample to determine the relative expression levels of a selected polynucleotide, particularly a polynucleotide corresponding to one or more of the differentially expressed genes described herein. Alternatively, gene expression in a test sample can be performed using serial analysis of gene expression (SAGE) methodology (*e.g.*, Velculescu et al., *Science* (1995) 270:484), differential display methodology (see, *e.g.*,
30 U.S. Patent Nos. 5,776,683 and 5,807,680), or DNA microarray methodology.

Alternatively, gene expression can be analyzed using hybridization analysis. Oligonucleotides or cDNA can be used to selectively identify or capture DNA or RNA of specific sequence composition, and the amount of RNA or cDNA hybridized to a known capture sequence determined qualitatively or quantitatively, to provide
5 information about the relative representation of a particular message within the pool of cellular messages in a sample.

Use of a single gene in diagnostic applications. The diagnostic methods of the invention can focus on the expression of a single differentially expressed gene represented by SEQ ID NO:1 or 3. For example, the diagnostic method can involve
10 detecting a differentially expressed gene, or a polymorphism of such a gene (*e.g.*, a polymorphism in a coding region or control region), that is associated with disease. Disease-associated polymorphisms can include deletion or truncation of the gene, mutations that alter expression level and/or affect binding specificity, such as interaction of the encoded protein with Notch receptor.

15 A number of methods are available for analyzing nucleic acids for the presence of a specific sequence, *e.g.*, a disease associated polymorphism. Where large amounts of DNA are available, genomic DNA is used directly. Alternatively, the region of interest is cloned into a suitable vector and grown in sufficient quantity for analysis. Cells that express a differentially expressed gene can be used as a source of
20 mRNA, which can be assayed directly or reverse transcribed into cDNA for analysis. The nucleic acid can be amplified by conventional techniques, such as the polymerase chain reaction (PCR), to provide sufficient amounts for analysis, and a detectable label can be included in the amplification reaction (*e.g.*, using a detectably labeled primer or detectably labeled oligonucleotides) to facilitate detection. Alternatively, various
25 methods are also known in the art that utilize oligonucleotide ligation as a means of detecting polymorphisms, see, *e.g.*, Riley et al., *Nucl. Acids Res.* (1990) 18:2887; and Delahunty et al., *Am. J. Hum. Genet.* (1996) 58:1239.

The amplified or cloned sample nucleic acid can be analyzed by one of a number of methods known in the art. The nucleic acid can be sequenced by dideoxy or
30 other methods, and the sequence of bases compared to a selected sequence, *e.g.*, to a

- wild-type sequence. Hybridization with the polymorphic or variant sequence can also be used to determine its presence in a sample (*e.g.*, by Southern blot, dot blot, *etc.*). The hybridization pattern of a polymorphic or variant sequence and a control sequence to an array of oligonucleotide probes immobilized on a solid support, as described in U.S. Patent No. 5,445,934, or in WO 95/35505, can also be used as a means of identifying polymorphic or variant sequences associated with disease. Single strand conformational polymorphism (SSCP) analysis, denaturing gradient gel electrophoresis (DGGE), and heteroduplex analysis in gel matrices are used to detect conformational changes created by DNA sequence variation as alterations in electrophoretic mobility.
- 10 Alternatively, where a polymorphism creates or destroys a recognition site for a restriction endonuclease, the sample is digested with that endonuclease, and the products size fractionated to determine whether the fragment was digested. Fractionation is performed by gel or capillary electrophoresis, particularly acrylamide or agarose gels.
- 15 Screening for mutations in a gene can be based on the functional or antigenic characteristics of the protein. Protein truncation assays are useful in detecting deletions that can affect the biological activity of the protein. Various immunoassays designed to detect polymorphisms in proteins can be used in screening. Where many diverse genetic mutations lead to a particular disease phenotype, functional protein
- 20 assays have proven to be effective screening tools. The activity of the encoded protein can be determined by comparison with the wild-type protein.

Use of Polypeptides to Screen for Peptide Analogs and Antagonists

- Polypeptides encoded by SEQ ID NOs:1 and 3 or by the corresponding full length genes can be used to screen peptide libraries to identify binding partners, such as ligands and receptors, from among the encoded polypeptides. Peptide libraries
- 25 can be synthesized according to methods known in the art (see, *e.g.*, U.S. Patent No. 5,010,175, and WO 91/17823). Screening methods may include, but are not limited to, yeast two-hybrid screens, protein co-immunoprecipitation assays, and lambda gt11 expression library screening. Agonists or antagonists of the polypeptides of the

invention can be screened using assay conditions that ideally resemble the conditions under which the native activity, *i.e.* interaction between Notch receptor and Notch ligand, is exhibited *in vivo*, that is, under physiologic pH, temperature, and ionic strength. Suitable agonists or antagonists will exhibit strong inhibition or enhancement
5 of the native activity at concentrations that do not cause toxic side effects in the subject. Agonists or antagonists that compete for binding to the native ligand can require concentrations equal to or greater than the native concentration, while inhibitors capable of binding irreversibly to the ligand can be added in concentrations on the order of the native concentration. In a non-limiting example, a peptide analog or antagonist would
10 be tested for its ability to affect interaction between a ligand of the invention and a Notch receptor.

Pharmaceutical Compositions and Therapeutic Uses

Pharmaceutical compositions of the invention can comprise polypeptides, antibodies, or polynucleotides (including antisense nucleotides and
15 ribozymes) of the claimed invention in a therapeutically effective amount. The term "therapeutically effective amount" as used herein refers to an amount of a therapeutic agent to treat, ameliorate, or prevent a desired disease or condition, or to exhibit a detectable therapeutic or preventative effect. The effect can be detected by, for example, chemical markers or antigen levels. Therapeutic effects also include reduction
20 in physical symptoms. The precise effective amount for a subject will depend upon the subject's size and health, the nature and extent of the condition, and the therapeutics or combination of therapeutics selected for administration. Thus, it is not useful to specify an exact effective amount in advance. However, the effective amount for a given situation is determined by routine experimentation and is within the judgment of the
25 clinician. For purposes of the present invention, an effective dose will generally be from about 0.01 mg/ kg to 50 mg/kg or 0.05 mg/kg to about 10 mg/kg of the polynucleotide constructs in the individual to which it is administered. A non-limiting example of a pharmaceutical composition is a composition that either enhances or diminishes binding of Notch to a ligand comprising a polypeptide of SEQ ID NO:2 and

4 or fragment thereof. Where the ligand/Notch interaction promotes a disease-related process, interference with the interaction would be the goal of the therapy. If normal ligand/Notch interaction does not occur because of a defect in one or the other protein, then use of a composition that restores normal function is appropriate.

5 In other instances it may be desirable to promote angiogenesis, for example in wound healing and in therapeutic angiogenesis, for example to treat ischemia.

A pharmaceutical composition can also contain a pharmaceutically acceptable carrier. The term "pharmaceutically acceptable carrier" refers to a carrier for
10 administration of a therapeutic agent, such as antibodies or a polypeptide, genes, and other therapeutic agents. The term refers to any pharmaceutical carrier that does not itself induce the production of antibodies harmful to the individual receiving the composition, and which can be administered without undue toxicity. Suitable carriers can be large, slowly metabolized macromolecules such as proteins, polysaccharides,
15 polylactic acids, polyglycolic acids, polymeric amino acids, amino acid copolymers, and inactive virus particles. Such carriers are well known to those of ordinary skill in the art. Pharmaceutically acceptable carriers in therapeutic compositions can include liquids such as water, saline, glycerol and ethanol. Auxiliary substances, such as wetting or emulsifying agents, pH buffering substances, and the like, can also be present
20 in such vehicles. Typically, the therapeutic compositions are prepared as injectables, either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid vehicles prior to injection can also be prepared. Liposomes are included within the definition of a pharmaceutically acceptable carrier. Pharmaceutically acceptable salts can also be present in the pharmaceutical
25 composition, *e.g.*, mineral acid salts such as hydrochlorides, hydrobromides, phosphates, sulfates, and the like; and the salts of organic acids such as acetates, propionates, malonates, benzoates, and the like. A thorough discussion of pharmaceutically acceptable excipients is available in *Remington's Pharmaceutical Sciences* (Mack Pub. Co., New Jersey, 1991).

Delivery Methods. Once formulated, the compositions of the invention can be (1) administered directly to the subject (*e.g.*, as polynucleotides or polypeptides); or (2) delivered *ex vivo*, to cells derived from the subject (*e.g.*, as in *ex vivo* gene therapy). Direct delivery of the compositions will generally be accomplished by
5 parenteral injection, *e.g.*, subcutaneously, intraperitoneally, intravenously or intramuscularly, myocardial, intratumoral, peritumoral, or to the interstitial space of a tissue. Other modes of administration include oral and pulmonary administration, suppositories, and transdermal applications, needles, and gene guns or hyposprays. Dosage treatment can be a single dose schedule or a multiple dose schedule.

10 Methods for the *ex vivo* delivery and reimplantation of transformed cells into a subject are known in the art and described in *e.g.*, International Publication No. WO 93/14778. Examples of cells useful in *ex vivo* applications include, for example, stem cells, particularly hematopoietic, lymph cells, macrophages, dendritic cells, or tumor cells. Generally, delivery of nucleic acids for both *ex vivo* and *in vitro*
15 applications can be accomplished by, for example, dextran-mediated transfection, calcium phosphate precipitation, polybrene mediated transfection, protoplast fusion, electroporation, encapsulation of the polynucleotide(s) in liposomes, direct microinjection of the DNA into nuclei, and viral-mediated, such as adenovirus or alphavirus, all well known in the art.

20 In a preferred embodiment, disorders of proliferation, such as tumor-associated angiogenesis, can be amenable to treatment by administration of a therapeutic agent based on the provided polynucleotide, corresponding polypeptide or other corresponding molecule (*e.g.*, antisense, ribozyme), agonist, or antagonist. The therapeutic agent can be administered in conjunction with one or more other agents
25 including but not limited to FGF, VEGF, angiopoietin, angiogenin, and thrombopoietin. Administered "in conjunction" includes administration at the same time, or within 1 day, 12 hours, 6 hours, one hour, or less than one hour, as the other therapeutic agent(s). The compositions may be mixed for co-administration, or may be administered separately by the same or different routes.

The dose and the means of administration of the inventive pharmaceutical compositions are determined based on the specific qualities of the therapeutic composition, the condition, age, and weight of the patient, the progression of the disease, and other relevant factors. For example, administration of polynucleotide therapeutic compositions agents of the invention includes local or systemic administration, including injection, oral administration, particle gun or catheterized administration, and topical administration. The therapeutic polynucleotide composition can contain an expression construct comprising a promoter operably linked to a polynucleotide of at least 12, 22, 25, 30, or 35 contiguous nucleotides of the polynucleotides disclosed herein. Various methods can be used to administer the therapeutic composition directly to a specific site in the body. For example, a small metastatic lesion is located and the therapeutic composition injected several times in several different locations within the body of tumor. Alternatively, arteries which serve a tumor are identified, and the therapeutic composition injected into such an artery, in order to deliver the composition directly into the tumor. A tumor that has a necrotic center is aspirated and the composition injected directly into the now empty center of the tumor. The antisense composition is directly administered to the surface of the tumor, for example, by topical application of the composition. X-ray imaging is used to assist in certain of the above delivery methods.

Receptor-mediated targeted delivery of therapeutic compositions containing an antisense polynucleotide, subgenomic polynucleotides, or antibodies to specific tissues can also be used. Receptor-mediated DNA delivery techniques are described in, for example, Findeis et al., *Trends Biotechnol.* (1993) 11:202; Chiou et al., *Gene Therapeutics: Methods And Applications Of Direct Gene Transfer* (J.A. Wolff, ed.) (1994); Wu et al., *J. Biol. Chem.* (1988) 263:621; Wu et al., *J. Biol. Chem.* (1994) 269:542; Zenke et al., *Proc. Natl. Acad. Sci. (USA)* (1990) 87:3655; Wu et al., *J. Biol. Chem.* (1991) 266:338. Therapeutic compositions containing a polynucleotide are administered in a range of about 100 ng to about 200 mg of DNA for local administration in a gene therapy protocol. Concentration ranges of about 500 ng to about 50 mg, about 1 mg to about 2 mg, about 5 mg to about 500 mg, and about 20 mg

to about 100 mg of DNA can also be used during a gene therapy protocol. Factors such as method of action (e.g., for enhancing or inhibiting levels of the encoded gene product) and efficacy of transformation and expression are considerations which will affect the dosage required for ultimate efficacy of the antisense subgenomic polynucleotides. Where greater expression is desired over a larger area of tissue, larger amounts of antisense subgenomic polynucleotides or the same amounts readministered in a successive protocol of administrations, or several administrations to different adjacent or close tissue portions of, for example, a tumor site, may be required to effect a positive therapeutic outcome. In all cases, routine experimentation in clinical trials will determine specific ranges for optimal therapeutic effect.

The present invention also relates to antisense oligonucleotides designed to interfere with the normal function of 3md3 and/or 2hd1 polynucleotides. Any modifications or variations of the antisense molecule which are known in the art to be broadly applicable to antisense technology are included within the scope of the invention. Such modifications include preparation of phosphorus-containing linkages as disclosed in U.S. Patents 5,536,821; 5,541,306; 5,550,111; 5,563,253; 5,571,799; 5,587,361, 5,625,050 and 5,958,773.

The antisense compounds of the invention can include modified bases as disclosed in 5,958,773 and patents disclosed therein. The antisense oligonucleotides of the invention can also be modified by chemically linking the oligonucleotide to one or more moieties or conjugates to enhance the activity, cellular distribution, or cellular uptake of the antisense oligonucleotide. Such moieties or conjugates include lipids such as cholesterol, cholic acid, thioether, aliphatic chains, phospholipids, polyamines, polyethylene glycol (PEG), palmityl moieties, and others as disclosed in, for example, U.S. Patents 5,514,758, 5,565,552, 5,567,810, 5,574,142, 5,585,481, 5,587,371, 5,597,696 and 5,958,773.

Chimeric antisense oligonucleotides are also within the scope of the invention, and can be prepared from the present inventive oligonucleotides using the methods described in, for example, U.S. Patents 5,013,830, 5,149,797, 5,403,711, 5,491,133, 5,565,350, 5,652,355, 5,700,922 and 5,958,773.

In the antisense art a certain degree of routine experimentation is required to select optimal antisense molecules for particular targets. To be effective, the antisense molecule preferably is targeted to an accessible, or exposed, portion of the target RNA molecule. Although in some cases information is available about the structure of target mRNA molecules, the current approach to inhibition using antisense is via experimentation. mRNA levels in the cell can be measured routinely in treated and control cells by reverse transcription of the mRNA and assaying the cDNA levels. The biological effect can be determined routinely by measuring cell growth or viability as is known in the art.

According to the invention, antisense oligonucleotides can be added to cultures of endothelial cells, and the effect on endothelial cell invasion can be determined, in order to identify antisense compositions useful for *in vivo* use.

Measuring the specificity of antisense activity by assaying and analyzing cDNA levels is an art-recognized method of validating antisense results. It has been suggested that RNA from treated and control cells should be reverse-transcribed and the resulting cDNA populations analyzed. (Branch, A. D., *T.I.B.S.* 23:45-50, 1998.)

The therapeutic polynucleotides and polypeptides of the present invention can be delivered using gene delivery vehicles. The gene delivery vehicle can be of viral or non-viral origin (see generally, Jolly, *Cancer Gene Therapy* (1994) 1:51; Kimura, *Human Gene Therapy* (1994) 5:845; Connelly, *Human Gene Therapy* (1995) 1:185; and Kaplitt, *Nature Genetics* (1994) 6:148). Expression of such coding sequences can be induced using endogenous mammalian or heterologous promoters. Expression of the coding sequence can be either constitutive or regulated.

Viral-based vectors for delivery of a desired polynucleotide and expression in a desired cell are well known in the art. Exemplary viral-based vehicles include, but are not limited to, recombinant retroviruses (see, *e.g.*, WO 90/07936; WO 94/03622; WO 93/25698; WO 93/25234; U.S. Patent No. 5, 219,740; WO 93/11230; WO 93/10218; U.S. Patent No. 4,777,127; GB Patent No. 2,200,651; EP 0 345 242; and WO 91/02805), alphavirus-based vectors (*e.g.*, Sindbis virus vectors, Semliki forest virus (ATCC VR-67; ATCC VR-1247), Ross River virus (ATCC VR-373; ATCC VR-

1246) and Venezuelan equine encephalitis virus (ATCC VR-923; ATCC VR-1250; ATCC VR 1249; ATCC VR-532), and adeno-associated virus (AAV) vectors (see, e.g., WO 94/12649, WO 93/03769; WO 93/19191; WO 94/28938; WO 95/11984 and WO 95/00655). Administration of DNA linked to killed adenovirus as described in Curiel, 5 *Hum. Gene Ther.* (1992) 3:147 can also be employed.

Non-viral delivery vehicles and methods can also be employed, including, but not limited to, polycationic condensed DNA linked or unlinked to killed adenovirus alone (see, e.g., Curiel, *Hum. Gene Ther.* (1992) 3:147); ligand-linked DNA (see, e.g., Wu, *J. Biol. Chem.* 264:16985 (1989)); eukaryotic cell delivery vehicles cells 10 (see, e.g., U.S. Patent No. 5,814,482; WO 95/07994; WO 96/17072; WO 95/30763; and WO 97/42338) and nucleic charge neutralization or fusion with cell membranes. Naked DNA can also be employed. Exemplary naked DNA introduction methods are described in WO 90/11092 and U.S. Patent No. 5,580,859. Liposomes that can act as gene delivery vehicles are described in U.S. Patent No. 5,422,120; WO 95/13796; WO 15 94/23697; WO 91/14445; and EP 0524968. Additional approaches are described in Philip, *Mol. Cell Biol.* 14:2411 (1994), and in Woffendin, *Proc. Natl. Acad. Sci.* (1994) 91:11581-11585.

Further non-viral delivery suitable for use includes mechanical delivery systems such as the approach described in Woffendin et al., *Proc. Natl. Acad. Sci. USA* 20 91(24):11581 (1994). Moreover, the coding sequence and the product of expression of such can be delivered through deposition of photopolymerized hydrogel materials or use of ionizing radiation (see, e.g., U.S. Patent No. 5,206,152 and WO 92/11033). Other conventional methods for gene delivery that can be used for delivery of the coding sequence include, for example, use of hand-held gene transfer particle gun (see, 25 e.g., U.S. Patent No. 5,149,655); use of ionizing radiation for activating transferred gene (see, e.g., U.S. Patent No. 5,206,152 and WO 92/11033).

The present invention will now be illustrated by reference to the following examples which set forth particularly advantageous embodiments. However, it should be noted that these embodiments are illustrative and are not to be construed as 30 restricting the invention in any way.

EXAMPLES

EXAMPLE 1

IDENTIFICATION AND ANALYSIS OF NOVEL NOTCH LIGANDS

5 Two ligands for the human Notch receptor have been identified, sequenced, and analyzed using Northern blots and *in situ* hybridization.

A. 3md3

 The first ligand is a 583 amino acid protein encoded by SEQ ID NO:1. The sequences are shown in Figure 5, and the ligand is referred to as 3md3.

10 On the basis of alignment with the mouse Delta 3 amino acid sequence, 3md3 is putatively identified as the human ortholog of mouse Delta 3 (Figure 1). Although a patent publication has identified a human "Delta 3" (WO 98 45434-A1), comparison of its alignment to mouse Delta 3, with 3md3 alignment to mouse Delta 3, indicates that 3md3 bears closer resemblance based on both amino acid (Figure 1) and
15 polynucleotide (Figure 3) comparisons.

 The expression of 3md3 in human tissues was investigated using an EST AI363919 (Image Clone No. 2016425) as a probe. The results from the Northern blot indicate that 3md3 was expressed as a 4.9 kb message in heart, kidney, skeletal muscle, and liver.

20 Table 1 shows the results of *in situ* hybridization of the 3md3 probe to normal and cancer tissue samples. Each result represents the data from six separate arrays. 3md3 is expressed at a higher level in the following types of cancer compared to the corresponding normal tissue: adrenal, lung, pancreas, and thyroid. 3md3 is expressed at a lower level in the following types of cancer compared to the
25 corresponding normal tissue: colon, esophagus, liver, prostate, and stomach. No expression was seen in samples from normal or cancerous breast, lymph node, and uterine tissue. In addition, 100% (3/3 samples, Table 2) of melanoma samples

expressed 3md3, suggesting its use as a marker and a potential therapeutic target in this type of cancer.

TABLE 1

In situ Expression of 3md3 in Normal and Cancer Tissue

Tissue Source	% expression in normal tissue	% expression in cancer tissue
Adrenal	0	33
Breast	0	0
Colon	24	8
Esophagus	50	43
Kidney	71	29
Liver	64	15
Lung	5	10
Lymph Node/ Lymphoma	0	0
Pancreas	55	75
Prostate	8	0
Stomach	25	6
Thyroid	0	33
Uterus	0	0

TABLE 2

In situ Expression of 3md3 in Cancer Tissue

Cancer: Tissue Source	% expression 3md3
Basal Cell	0
Cervical	33
Chorio	33
Epithelial	0
Fibro	0
Germ Cell	0
Leiomyo	0
Melanoma	100
Seminoma	33

B. 2hd1

The second ligand is a protein encoded by SEQ ID NO:3. The sequences
5 are shown in Figure 6, and the ligand is referred to as 2hd1.

EXAMPLE 2

EFFECT OF ANTISENSE OLIGONUCLEOTIDES ON ENDOTHELIAL CELL INVASION

Adrenal cortex-derived bovine microvascular endothelial (BME) cells
10 are grown in an α -modified minimal essential medium (Life Technologies, Inc. AG, Basel, Switzerland), supplemented with 15% heat-inactivated donor calf serum (DCS, Life Technologies, Inc.), penicillin (110 units/ml), and streptomycin (110 μ g/ml). BME cells are subcultured at a 1:4 split ratio in 1.5% gelatin-coated tissue culture dishes or flasks (Falcon Labware, Becton Dickinson Company, Lincoln Park, NJ). The *in vitro*
15 angiogenesis assay is performed as described (Montesano, R. et al., *Cell* 42:469-477, 1985) in 16-mm tissue culture wells (Nunc, A/S Nunc, Roskilde, Denmark). BME cells are seeded at $5.0-7.5 \times 10^4$ cells/well in 500 μ l of α -modified minimal essential

medium, 5% DCS (donor calf serum). Prior to reaching confluence (less than 3 days), DCS is further reduced to 2%, and cells are treated with recombinant human FGF-2, recombinant human VEGF, and antisense oligonucleotides. Oligonucleotides are added to the cells 2 h before cytokines on the first day of treatment. Medium and cytokines
5 are renewed after 2 days, and oligonucleotides are added either every day or every other day during the 4-day assay period. Cultures are fixed *in situ* after a further 2 days with 2.5% glutaraldehyde in 100 mM sodium cacodylate buffer (pH 7.4), and photographed. For quantitation, randomly selected fields measuring 1.0 x 1.4 mm are photographed in each well at a single level beneath the surface monolayer by phase contrast microscopy,
10 using a Nikon Diaphot TMD inverted photomicroscope. In each experiment, invasion is quantitated from at least three photographic fields by determining the total additive length of all cellular structures that penetrate beneath the surface monolayer either as apparently single cells or in the form of cell cords or tubes. (Pepper, M.S. et al., *Biochem. Biophys. Res. Commun.* 189:824-831, 1992.)

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EXAMPLE 3

EXPRESSION OF 3MD3 IN HUMAN MICROVASCULAR ENDOTHELIAL CELLS

The expression of 3md3 in human microvascular endothelial cells (HMEC) is measured by assaying the mRNA in cells after a variety of treatments.
20 mRNA expression levels are assayed by preparing probes for Northern blots as described above. bFGF-treated HMEC are prepared by incubation with bFGF at 10 mg/ml for 2 hours. VEGF-treated HMEC are prepared by incubation with 20 ng/ml VEGF for 2 hours. Following incubation with the respective growth factor, the cells are washed and lysis buffer added for RNA preparation.

25

The HMEC treated as above can also be exposed to one or more antisense oligomers based on SEQ ID NO:1 or 3, using methods described by Zimrin, A. et al., *J. Biol. Chem.* 271:32499-32502, 1996.

EXAMPLE 4

EFFECT OF 3MD3 EXPRESSION ON TUMOR INDUCED ANGIOGENESIS *IN VIVO*

The effect of 3md3 on tumor growth and tumor induced angiogenesis is
5 assessed by:

(a) Infecting the highly metastatic human breast cancer cell line
MDA-MB-231 with retroviruses encoding (i) the extracellular domain of 3md3,
and (ii) full length 3md3. The infected cells are subsequently implanted in nude
mice and the growth of the tumors as compared to control groups is monitored
10 over time, and

(b) First establishing a tumor mass by implanting MDA-MB-231
into nude mice. Adenoviruses encoding (i) the extracellular domain of 3md3,
and (ii) full length 3md3 are then injected intratumorally and peritumorally. The
growth of the tumors as compared to control groups is monitored over time.

15

EXAMPLE 5

EFFECT OF 3MD3 ON VEGF AND bFGF INDUCED ANGIOGENESIS *IN VIVO*

Adenoviruses encoding (i) the extracellular domain of 3md3, and (ii) full
length 3md3 are injected into the retroperitoneal adipose tissue alone or in combination
20 with adenoviruses encoding VEGF or bFGF using methods described by Magovern,
C.J. et al., *Hum. Gene Ther.* 8:215, 1997, and U.S. Patent No. 5,869,037.
Neovascularization of the adipose tissue is assessed on day 20 and day 30 post-injection
by histological evaluation.

25

From the foregoing it will be appreciated that, although specific
embodiments of the invention have been described herein for purposes of illustration,
various modifications may be made without deviating from the spirit and scope of the
invention. Accordingly, the invention is not limited except as by the appended claims.

CLAIMS

WE CLAIM:

1. An isolated nucleic acid molecule comprising a polynucleotide selected from the group consisting of:
 - (a) a polynucleotide encoding amino acids from about 1 to about 583 of SEQ ID NO:2;
 - (b) a polynucleotide encoding amino acids from about 2 to about 583 of SEQ ID NO:2;
 - (c) a polynucleotide encoding amino acids from about 1 to about 81 of SEQ ID NO:4;
 - (d) a polynucleotide encoding amino acids from about 2 to about 81 of SEQ ID NO:4;
 - (e) the polynucleotide complement of the polynucleotide of (a)-(d);and
 - (f) a polynucleotide at least 90% identical to the polynucleotide of (a)-(e).
2. An isolated nucleic acid molecule consisting of a nucleic acid comprising 50-1752 contiguous nucleotides from the coding region of SEQ ID NO:1.
3. The isolated nucleic acid molecule of claim 2, which comprises 100-1500 contiguous nucleotides.
4. The isolated nucleic acid molecule of claim 3, which comprises 500-1000 contiguous nucleotides.
5. An isolated nucleic acid molecule comprising a polynucleotide encoding a polypeptide wherein, except for at least one conservative amino acid

substitution, said polypeptide has an amino acid sequence selected from the group consisting of:

- (a) amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) amino acids from about 2 to about 583 of SEQ ID NO:2;
- (c) amino acids from about 1 to about 81 of SEQ ID NO:4; and
- (d) amino acids from about 2 to about 81 of SEQ ID NO:4.

6. The isolated nucleic acid molecule of claim 1, which is DNA.

7. A method of making a recombinant vector comprising inserting a nucleic acid molecule of claim 1 into a vector in operable linkage to a promoter.

8. A recombinant vector produced by the method of claim 7.

9. A method of making a recombinant host cell comprising introducing the recombinant vector of claim 8 into a host cell.

10. A recombinant host cell produced by the method of claim 9.

11. A recombinant method of producing a polypeptide, comprising culturing the recombinant host cell of claim 10 under conditions such that said polypeptide is expressed and recovering said polypeptide.

12. An isolated polypeptide comprising amino acids at least 95% identical to amino acids selected from the group consisting of:

- (a) amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) amino acids from about 2 to about 583 of SEQ ID NO:2;
- (c) amino acids from about 1 to about 81 of SEQ ID NO:4; and
- (d) amino acids from about 2 to about 81 of SEQ ID NO:4.

13. An isolated polypeptide wherein, expect for at least one conservative amino acid substitution, said polypeptide has an amino acid sequence selected from the group consisting of:

- (a) amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) amino acids from about 2 to about 583 of SEQ ID NO:2;
- (c) amino acids from about 1 to about 81 of SEQ ID NO:4; and
- (d) amino acids from about 2 to about 81 of SEQ ID NO:4.

14. An isolated polypeptide comprising amino acids selected from the group consisting of:

- (a) amino acids from about 1 to about 583 of SEQ ID NO:2;
- (b) amino acids from about 2 to about 583 of SEQ ID NO:2;
- (c) amino acids from about 1 to about 81 of SEQ ID NO:4; and
- (d) amino acids from about 2 to about 81 of SEQ ID NO:4.

15. An epitope-bearing portion of a polypeptide comprising consisting of SEQ ID NO:2.

16. The epitope-bearing portion of claim 15, which comprises about 5 to about 50 contiguous amino acids.

17. The epitope-bearing portion of claim 16, which comprises about 10 to about 20 contiguous amino acids.

18. An isolated antibody that binds to the polypeptide of claim 12.

19. An isolated antibody that binds to the polypeptide of claim 13.

20. An isolated antibody that binds to the polypeptide of claim 14.

21. A complex comprising a protein comprising an amino acid sequence selected from the group consisting of SEQ ID NO:2 and SEQ ID NO:4.

22. A complex comprising a fragment of the amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and a Dishevelled protein wherein said fragment is capable of forming a complex with said Dishevelled protein.

23. A pharmaceutical composition comprising a therapeutically effective amount of a polypeptide selected from the group consisting of SEQ ID NO:2 and SEQ ID NO:4, and a pharmaceutically effective carrier.

24. A method of detecting Notch ligand expression in human cancer cells, said method comprising:

obtaining mRNA from said cells; and

contacting said mRNA with a polynucleotide of SEQ ID NO:1 under stringent hybridization conditions, wherein formation of a duplex comprising a polynucleotide of SEQ ID NO:1 indicates expression of Notch ligand wherein said Notch ligand is encoded by a gene comprising SEQ ID NO:1 or its complement.

25. A method of detecting Notch ligand expression in human melanoma cells, said method comprising:

obtaining mRNA from said cells; and

contacting said mRNA with a polynucleotide of SEQ ID NO:1 under stringent hybridization conditions, wherein formation of a duplex comprising a polynucleotide of SEQ ID NO:1 indicates expression of Notch ligand wherein said Notch ligand is encoded by a gene comprising SEQ ID NO:1 or its complement.

26. A method of enhancing angiogenesis in a mammal in need thereof, said method comprising administering the composition of claim 23 and at least one growth factor selected from the group consisting of bFGF and VEGF.

27. The method of claim 26 wherein said mammal exhibits tissue ischemia.

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3dm3p      .....M VSRMSGLLS QTVIDALIFL PQTTPAGVFE LQIHSGGPGP QPGAPSPCS ARL.....PC RLFRVCLKP GLSEEAESP DALGAAL SAR GMYT 91
moused113p checksun.M VSLQVSP.LS QTLILAF.LL PQALPAGVFE LQIHSGGPGP QLGTPSPSPCN ARG.....PC RLFRVCLKP GVSQEAESL CALGAALSTS VPMYT 97
hdelta1p  .....M GSRCALALAV LSALLCOVW. ....SSGVFE LKLEDFVNNK QLLGNCCR GGAGPPCAC RTIFRVCLKH YQASVSPDP QTVGSANTPV LQVDS 91
w80813     .....maa asrsasgwa1 1..lljvalwq qraagsgvfj lqldfiner gv1..... asgrpcepge rtf frcvclkh fqawvspg p dtfgt vstopv lgtus 88

3dm3p      EQPGAPAPDL PLPDGLLQVPI FRDAMPGTFS FLIETMREEL GQDIGGPW. ...SLLARVAG RRRLAAGGPW ARDTPAGAW ETRQSYRARC EPPAVGTACT RLCRP 193
moused113p EHPGESAAAL PLPDGLVRVP FRDAMPGTFS FLIETMREQL GEHAGGPW. ...NLLARVVG RRRLAAGGPW ARDTPAGAW ETRQSYRARC EPPAVGAACA RLCRS 199
hdelta1p  FSLPDGGGAD SAFSNPIRFP FGFIMPGTFS LTIETALHTDS PDLATEN. PERLISB IAT QRLHIVGEEM SQDLHSSGRT DLKYSYRVC DEHYVGECS VFCRP 194
w80813     favrddssgg g..rnp1qlp fnftwpatfs l1ieaaha.p gddlrpealp pda1lisk1ai qgs1avqnmw 1ldegt1t r1rysyrv1b sdryvgdncs r1ckk 190

3dm3p      RSAP.....S RCGPQLRPCA PLEDE.....S VORAGCSPEH GFDEQPGECR QLEGMTP1CQ TVPVSTSSCL SP..... RBPSS 261
moused113p RSAP.....S RCGPQLRPCT PFPDECAPS VORPGCSPEH GYDEPDECR QLEGMTP1CQ TVPVSTSSCL NS..... RMPJP 217
hdelta1p  RDAFGHFTC GERGEKVCNP GWKGPVCTEP IQLPGDEQH GFCDKPECK QRVGMQCRYC DECIRYPGCL HGTCQQPWQC NCQEGWGLF CNQDLNYCTH HKPCK 299
w80813     hndhfhgryc qpdgn1lsc1p gwtdeycqqp id1sghegn gyaskpaec1 qrpwqgr1l1 neciphngtr lgtcstpwc tcdegwglf cdqdlnycth hskk 295

3dm3p      ATTGLVP3P QP..... CDGIPCAN GGSQSETPRS FEQTCPRGFY GLRCEVSGVIT CADGPCFNGG LQVGADPDS AMIQCPPGF QGSNQ 346
moused113p ASTGLLP3P QP..... CDGIPCAN GGSQSETSGS FEQACPRGFY GLRCEVSGVIT CADGPCFNGG LQVGEDPDS XVMQCPPGF QGSNQ 356
hdelta1p  NGATQNTN1GQ QSYTCSCRPG YTGATCELGI DECDSYCKN GGSQTDLENS YSQTCPPGFY GKICELSAMT CADGPCFNGG RQSDSPJ.GG YSQCPCVGY SQFNG 402
w80813     ngatbsnsq rsytcrcrp ytgvdcele1 secdsnpctn ggsckdqdg ytd1cpdw g1jncst1s1 cadspcfnng sgrernq.ga ryacecpphf tgsng 399

3dm3p      EKRVDRCSLQ PCNIGGLCLD LQHALRCRCR AGFAGPRCEH D1DDCAGRAC ANGCTVEGG GAHRCSCALG FGGRDCRERA ...DPCAARPC AHGRCYAHF S..QL 447
moused113p EKRVDRCSLQ PCQNGGLCLD LGHAXXRCR AGFAGPRCEH D1DDCAGRAC ANGCTVEGG GSRRCSCALG FGGRDCRERA ...DPCAARPC AHGRCYAHF S..QL 457
hdelta1p  EKK1DYCSS PCSNGAKCVD LGDAYLRCRQ AGFSRHCDD NVDDCASSPC ANGCTORDG NDFSC1CPPG YTGRCSPV S..RDEHAPC HNGATCHE...RGHY 503
w80813     ekk1vdrctsn pdanggc1h rpsrmrcr1 pgrf1gt1cel hvscarnpc1 ahgtgchdle ng1ngt1cpag1 f1sgrrcevft1 s1daccasspc1 fnratcyt1d11st1dtf 504

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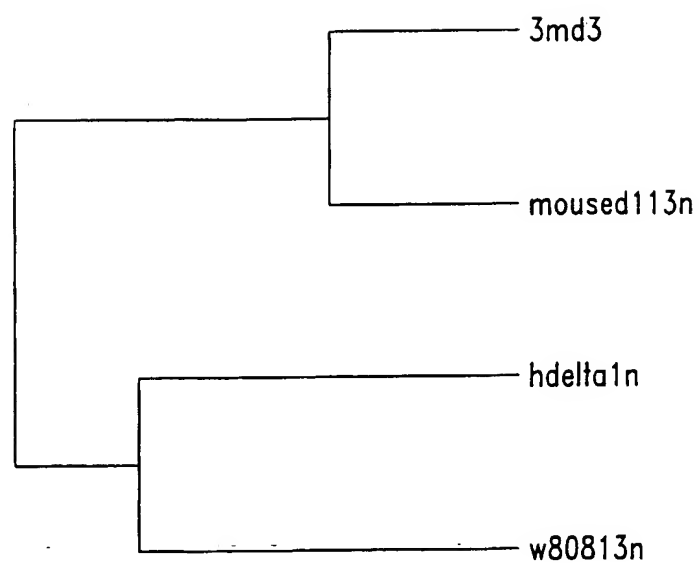
Fig. 1A

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3dm3p	VOACAPGYVG	ARCEFPV	HPDGASALPA	APPGLRPGDP	QRYLLPPALG	LLVAAGVAGA	ALLMIVRRR	GHSQDAGSRL	LAGTPEPSVH	ALPDALNNL	543	
moused113p	VOACAPGYVG	VRCEFFM	RPDGADAVPA	APRGLRQADP	QRFLPPALG	LLVAAGVAGA	ALLIVIVRRR	GPGQDTGTRL	LSGTREPSVH	TLPDALNNL	553	
hdel1alp	VOECAPGYGG	PNCQFLPEL	PPGPVAVDLT	EKLEGGGPF	PWAVCAGVI	LLMLLLGC	AAVMQVRLR	LOKHR	PPADPCR	GETETMNLIA	599	
w80813	vgncpygfvg	srcefpv	g1	pps	pwavslgv	llayllvllg	llmavavrl	qlrlr	mpddgsr	edhmls	577	
3dm3p	...	RTQEGS	GDGPSSSDW	NRPEVDPPQ	IYVISAPSIY	AREA	583	
moused113p	...	RLQDGA	GDGPSSSDW	NHPEDGDSRS	IYVIPAPSIY	AREA	593	
hdel1alp	KDISVSIIGA	TQIKNTNKKA	DFHGDHSADK	NGFKARYPA	VDYNLVQDLK	GDITAVRDAH	SKRDTKCQDQ	GSSGEEKGTP	TTLRGEASE	RKRPDGSGCST	703	
w80813	n	lipa	aqlkntnqkk	elevdcgldk	sncgkqnh	ldynladgp	grgt	mpgk fphs	dkslgeka.p	lrhsekpec r	isames prdsm	665
3dm3p	583	
moused113p	593	
hdel1alp	YQSVYVISEE	KOECVIATEV	723	723	
w80813	yqsvclisee	rncviatev	685	685	

Fig. 1B

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*Fig. 2*

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3md3	ATGGTCTCTCC	CACGATGTC	GGGGTCTCTG	TGGGATGTC	TGATCTTAC	GCTCATTTTC	CTTCCCTCAGA	CAACGCTCCG	TGGGCTCTTC	GAGCTGCAGA	TCCAC	105
moused113n	ATGGTCTCTC	TGCAGTCTC	...TCCGCTT	TGGGATGTC	TGATCTTAC	...TTTCTTT	CTTCCCTCAGA	CACTGCAGC	TGGGCTCTTC	GAGCTGCAGA	TTTCA	99
hdel1aln	ATGGG	TGGGATGTC	TGATCTTAC	CTGGGCTTTC	CTTCCCTCAGA	TCTGCAGC	TGGGCTCTTC	GAGCTGCAGA	TGCAG	90
w80813n	ATGGGCTCTC	CGTCCCGAG	CGCCTCTGG	TGGGATGTC	TGATCTTAC	GGGCTTTC	CTTCCCTCAGA	GGGCTCTTC	TGGGCTCTTC	GAGCTGCAGA	TGCAG	105
3md3	TCCTTCGGGC	CGGCTCAGG	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	195
moused113n	TCCTTCGGGC	CAGGCTCAGG	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	189
hdel1aln	GAGTTCTGTC	ACAAGAAGGG	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	195
w80813n	GAGTTCTGTC	ACAAGAAGGG	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	189
3md3	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	300
moused113n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	294
hdel1aln	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	300
w80813n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	289
3md3	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	405
moused113n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	399
hdel1aln	GACTTGGGCT	TCAGGAGGCC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	405
w80813n	GACTTGGGCT	TCAGGAGGCC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	387
3md3	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	501
moused113n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	495
hdel1aln	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	504
w80813n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	492
3md3	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	606
moused113n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	600
hdel1aln	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	609
w80813n	CTCTGGGCTCT	GAGAGGAGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	CCCTTCGGGC	597

Fig. 3A

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3md3	684	GTGAA
moused113n	690	GTGAA
hdel1a1n	714	GTGAC
w80813n	702	ATCAYC
3md3	765
moused113n	771
hdel1a1n	819	GGCAG
w80813n	807	ggcaa
3md3	804
moused113n	810
hdel1a1n	924	CGGGC
w80813n	912	gtggg
3md3	855	GCIGI
moused113n	861	GCIGI
hdel1a1n	1029	GCIGI
w80813n	1017	gctgt
3md3	960	AGGAC
moused113n	966	ATGGC
hdel1a1n	1134	AGGAC
w80813n	1122	abggg
3md3	1065	GGCTG
moused113n	1071	GGCTG
hdel1a1n	1233	GGCTG
w80813n	1224	qbagc

Fig. 3B

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3md3	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1170
moused113n	GAACCAATGCT	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	GTGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1176
hdel1aln	TCACCGCTG	CTATATGCTG	CAAGTATGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	GTGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1338
w80813n	adacccctgtg	cdagaggggg	adagtgccctg	adccgaggtg	caagcgcat	gtgcgcgctg	cgctcdggat	tgacgggac	ctactgtgah	tdtdacgltka	dcgac	1329
3md3	TGCGCGCGCG	GGGCTGCGCG	TAACGGCGCG	AGTATGCTG	AGCGCGCGCG	GTGCGCGCTG	TGCGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1269
moused113n	TGCGCGCGCG	GGGCTGCTG	TAACGGCGCG	AGTATGCTG	AGCGCGCGCG	GTGCGCGCTG	TGCGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1275
hdel1aln	TGCGCGCTG	GGGCTGCGCG	TAACGGCGCG	AGTATGCTG	AGCGCGCGCG	GTGCGCGCTG	TGCGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1437
w80813n	tgbgccclta	adcdltgcgc	ddacggltggc	adltggcatb	acctggagaa	tggggtcatg	tggaacctgc	dtgcccggctb	gtctggccgh	cgclgtgag	tgcg	1434
3md3	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1308
moused113n	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1374
hdel1aln	GCACCGCTG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1536
w80813n	acatccatcg	atgctgtgtg	tdtbgagltcc	tggttcacaa	gggcccacctg	ctacacgag	dtctccacag	atgctgtgtg	gtggaactgc	ddtaltggct	tlgtg	1539
3md3	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1473
moused113n	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1479
hdel1aln	GCACCGCTG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1640
w80813n	ggcagcctct	gcgaggttcc	dgtgggcttg	ccgcccccccccc	1592
3md3	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1578
moused113n	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1584
hdel1aln	GCACCGCTG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1745
w80813n	cdtctgtgtg	gdtgtggggg	tgccagtgtct	gcctgtatctgccccccc	1688
3md3	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1683
moused113n	GAACCAATGCG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1689
hdel1aln	GCACCGCTG	GAATATGCGCG	AGTATGCGCTG	GACCTATGCGCG	AGCGCTGCG	GTGCGCGCTG	CGCGCGCGCG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	TGCGCGCGCTG	1847
w80813n	cagc.....	aggggag	ccatgaaacaa	dtgttctgtgccccccc	1766

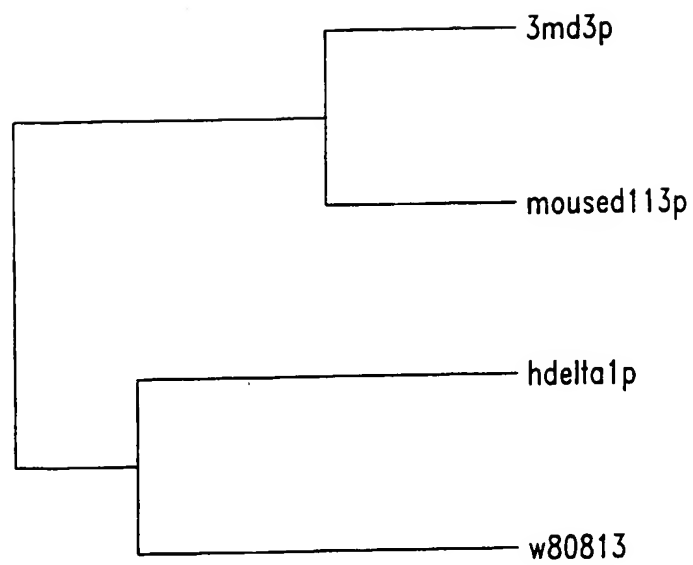
Fig. 3C

3md3	CTGTGAAGATG	TAGACCTTCA	AGGCAATTTAT	GTGATATCTG	CTGCTTCCAT	CTAGGCTTGG	GTAGGCTTGA	1752
moused113n	CGTGAAGATG	GAGACTCTAG	ATCCATTTAT	GTGATACCTG	CCGCTTCCAT	TTATGCAACA	GAGGCTTGA	1758
hdel1aln	GAAGAAGGCG	GACTTCTACG	GGGACCAACG	CGCCGACAAAG	...AATGGCT	TTAAGGCTCG	CTACCCGAGCG	1949
w80813n	cdagaagagag	gagctggaag	tggactcttgg	ccctggagacag	tggactctg	gdaaactagca	aaaddacaca	1868
3md3	1752
moused113n	1758
hdel1aln	CGACACCGCC	GTCAGGGACG	CGCACAGCAA	GGGTGACACC	AAGTGCACG	CCCAGGGCTC	CTCAGGGGAG	2054
w80813n	gcgggggacc	atgccaggaa	agtttccca	cagtgacaag	agcttaggag	agaaggcgcc	1954
3md3	1752
moused113n	1758
hdel1aln	AGCATCTGAA	AGAAAAGGC	CGGACTCGGG	CTGTTCACCT	TCAAAAGACA	CCAAGTACCA	GTCGGTGTAC	2159
w80813n	agtgct	ggatatcagc	gatgtgtctcc	cccaggact	ccatgtacca	2045
3md3	1752
moused113n	1758
hdel1aln	AACTGAGGTG	TAAATGGAA	GTGA	2183				
w80813n	cacggaggta	2055				

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Fig. 3D

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*Fig. 4*

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3md3 Nucleotide Sequence: 1752 (SEQ ID NO:1)

ATGGTCTCCCCACGGATGTCCGGGCTCCTCTCCCAGACTGTGATCCTAGC
GCTCATTTTCTCCCCAGACACGGCCCGCTGGCGTCTTCGAGCTGCAGA
TCCACTCTTTCGGGCCGGGTCCAGGCCCTGGGGCCCCGCGGTCCCCCTGC
AGCGCCCGGCTCCCCTGCCGCCTCTTCTTCAGAGTCTGCCTGAAGCCTGG
GCTCTCAGAGGAGGCCGCCGAGTCCCCGTGCGCCCTGGGCGGGCGCTGA
GTGCGCGCGGACCGGTCTACACCGAGCAGCCCGAGCGCCCGCGCCTGAT
CTCCCACTGCCCGACGGCCTCTTGAGGTGCCCTTCCGGGACGCCTGGCC
TGGCACCTTCTCTTTCATCATCGAAACCTGGAGAGAGGAGTTAGGAGACC
AGATTGGAGGGCCCCGCTGGAGCCTGCTGGCGCGCTGGCTGGCAGGCGG
CGCTTGGCAGCCGGAGGCCCGTGGGCCCGGGACATTCAGCGCGCAGGCGC
CTGGGAGCTGCGCTGCTCGTACCGCGCGCGCTGCGAGCCGCTGCGGTGCG
GGACCGCGTGACGCGCCTTGCCGTCCGCGCAGCGCCCCCTCGCGGTGC
GGTCCGGGACTGCGCCCCCTGCGCACCGCTCGAGGACGAATCGGTGTGCCG
AGCAGGCTGCAGCCCTGAGCATGGCTTCTGTGAACAGCCCGGTGAATGCC
GATGCCTAGAGGGCTGGACTGGACCCCTCTGCACGGTCCCTGTCTCCACC
AGCAGCTGCCTCAGCCCCAGGGGCCCGTCTCTGCTACCACCGGATGCCT
TGTCCTGGGCCCTGGGCCCTGTGACGGGAACCCGTGTGCCAATGGAGGCA
GCTGTAGTGAGACACCCAGGTCCTTTGAATGCACCTGCCCGCGTGGGTTT
TACGGGCTGCGGTGTGAGGTGAGCGGGGTGACATGTGCAGATGGACCCTG
CTTCAACGGCGGGCTTGTGTGTGCGGGGTGCAGACCCTGACTCTGCCTACA
TCTGCCACTGCCCACCTGGTTTTCAAGGCTCCAAGTGTGAGAAGAGGGTG
GACCGGTGCAGCCTGCAGCCATGCCGCAATGGCGGACTCTGCCTGGACCT
GGGCCACGCCCTGCGCTGCCGCTGCCGCGCCGGCTTCGCGGGTCTCGCT
GCGAGCACGACCTGGACGACTGCGCGGGCCGCGCCTGCGCTAACGGCGGC
ACGTGTGTGGAGGGCGGCGGCGCGCACCGCTGCTCCTGCGCGCTGGGCTT
CGGCGGGCCGCGACTGCCGCGAGCGCGCGGACCCGTGCGCCGCGCGCCCCT
GTGCTACGGCGGGCGCTGCTACGCCCACTTCTCCGGCCTCGTCTGCGCT
TGCCTCCCGGTACATGGGAGCGCGGTGTGAGTTCCAGTGCACCCCGA
CGGCGCAAGCGCCTTGCCCGCGGCCCGCGGGCCTCAGGCCCGGGGACC
CTCAGCGCTACCTTTTGCTCCGGCTCTGGGACTGCTCGTGGCCGCGGGC
GTGGCCGGCGCTGCGCTCTTGCTGGTCCACGTGCGCCGCGGTGGCCACTC
CCAGGATGCTGGGTCTCGCTTGCTGGCTGGGACCCCGGAGCCGTGAGTCC
ACGCACTCCCGGATGCACTCAACAACCTAAGGACGCAGGAGGGTTCCGGG
GATGGTCCGAGCTCGTCCGTAGATTGGAATCGCCCTGAAGATGTAGACCC
TCAAGGGATTTATGTATATCTGCTCCTTCATCTACGCTCGGGAGGCCT
GA

Fig. 5A

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3md3 Protein Sequence: 583 (SEQ ID NO: 2)

MVSPRMSGLLSQTIVILALIFLPQTRPAGVFELQIHSFGPGPGPGAPRSPC
SARLPCRLFFRVCLKPGLSEEAESPALGAALSARGPVYTEQPGAPAPD
LPLPDGLLQVPFRDAWPGTFSFIIETWREELGDQIGGPAWSLLARVAGRR
RLAAGGPWARDIQRAGAWELRCSYRARCEPPAVGTACTRLCRPRSAPSRC
GPGLRPCAPLEDESVCRAGCSPEHGFCEQPGEERCLEGTGPLCTVPVST
SSCLSPRGSSATTGCLVPGPGPCDGNPCANGGSCSETPRSFECTCPRGF
YGLRCEVSGVTCADGPCFNGGLCVGGADPDSAYICHCPPGFQGSNCEKRV
DRCSLQPCRNGGLCLDLGHALRCRCRAGFAGPRCEHLDLDCAGRACANGG
TCVEGGGAHRCSCALGFGGRDCRERADPCAARPCAHGGRCYAHFSGLVCA
CAPGYMGARCEFPVHPDGASALPAAPPGLRPGDPQRYLLPPALGLLVAAG
VAGAALLLVHVRRRGHSQDAGSRLLAGTPEPSVHALPDALNNLRTQEGSG
DGPSSSV DWNRPEDVDPOGIYVISAPSIYAREA

Fig. 5B

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2hd1 Nucleotide Sequence: 1307 (SEQ ID NO: 3)

AGTACTCCTACCGCTTCGTGTGTGACGAACACTACTACGGAGAGGGCTGCTCCGTTTTCT
GCCGTCCCCGGGACGATGCCTTCGGCCACTTCACCTGTGGGGAGCGTGGGGAGAAAGTGT
GCAACCCTGGCTGGAAAGGGCCCTACTGCACAGAGCGTGAGTCTCTGGGAAGGCACCGCT
GGCTCACTCGTCCACGAACACGGACCACGCGCAGGGACGGGGCTTCCTGAGCCACGGGGG
GCTTGGGACTGTAGAGATGTTCTGGTGGGGAACTGAGGCCAGAGGACAGAAGTGGATT
GCTATAAGTCACAGCTCGTCAGTGGGGGGGTGGGGTCAACGCAGACATTTTAACATCCC
AGGCTGTGTTTATCCACTATCGGAAGTGCCTTTCTTAATCAGGGAGGATTTTAGAGACAG
GGCCAGGGGTGAGGAAGTAAAGCCAGTGCTACCCCAAGGGTGTGTGATTAGAGAGGGAG
AGGAGGAAGGAAGGGAGGAACACAGAGAGAGCTTGTGTGTCAGGGGCACCATTTCAACCC
GAGTTCCAGTGCTGGAACAGCATCACACTGGGAAACGTTCCATTTTCTCTCTGGAGCTG
GTGTGCTTGACCTCTCTGGAGCAAACGCCTTCCGGATACTCCCTGTGACACGCACTGTC
TATGCTGGCCAGAGAGCAGGCTTTCACCTCTGTGGGCTGCTGAGGCCAGGTCTCCAAGGC
CTGTGTGGGCGAGGGGTGCACAGCCCCGTCTGGCTTGAATGCTCAGGCAGCACCTTGTCT
GGAAAAGCAATGTCTTCCCAATAGTGACAGAGGCTCTACCTGCCTCTTATTAGGTATTGA
TGTGTCAATGTCATGGCAGGCAGGTGACTAGGGCAGGGTGGGGCCGTGCTGGCTCCTGG
TTCTGGCTCATGGGGACCTCAGGAGCCCTCTCTCCAGCTGACTGAGGCCTCGCCTGCACG
CCTGGCCGTCCCAGCCCATTGGTACCGGATTTCTCTACAGCTGGGGATTGGGTAGGTCCT
GGAGCTGCCAGAACTCCAGGGAAGTGCATTCTCCTTCCCTGGAACTGGACAACCTTG
GAGAGGGGCTCTGGGAGGCCCAGAACCTCTGGCAGGAGCTGGGTAGTGCCTGGGGTTGAG
GGTGGGTCTTCCCATTCAGTGAGTGCCTTGATGTCCTTGCTCCTTAGCTTCCCAAATTCC
CTCCGGAAGTTACTGAGCTCCTTCTAAGCTTTGCCTTGGCCTGAACTGGTTCTGGGGAAA
AA

2hd1 Protein Sequence: 81 (SEQ ID NO: 4)

GRTDLKYSYRFVCDHYHGGCSVFCRPRDDAFGHFTCGERGEKVCNPGWKGPYCTERES
LGRHRWLTRPRTRTTRDGAS

Fig. 6

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Notch signaling

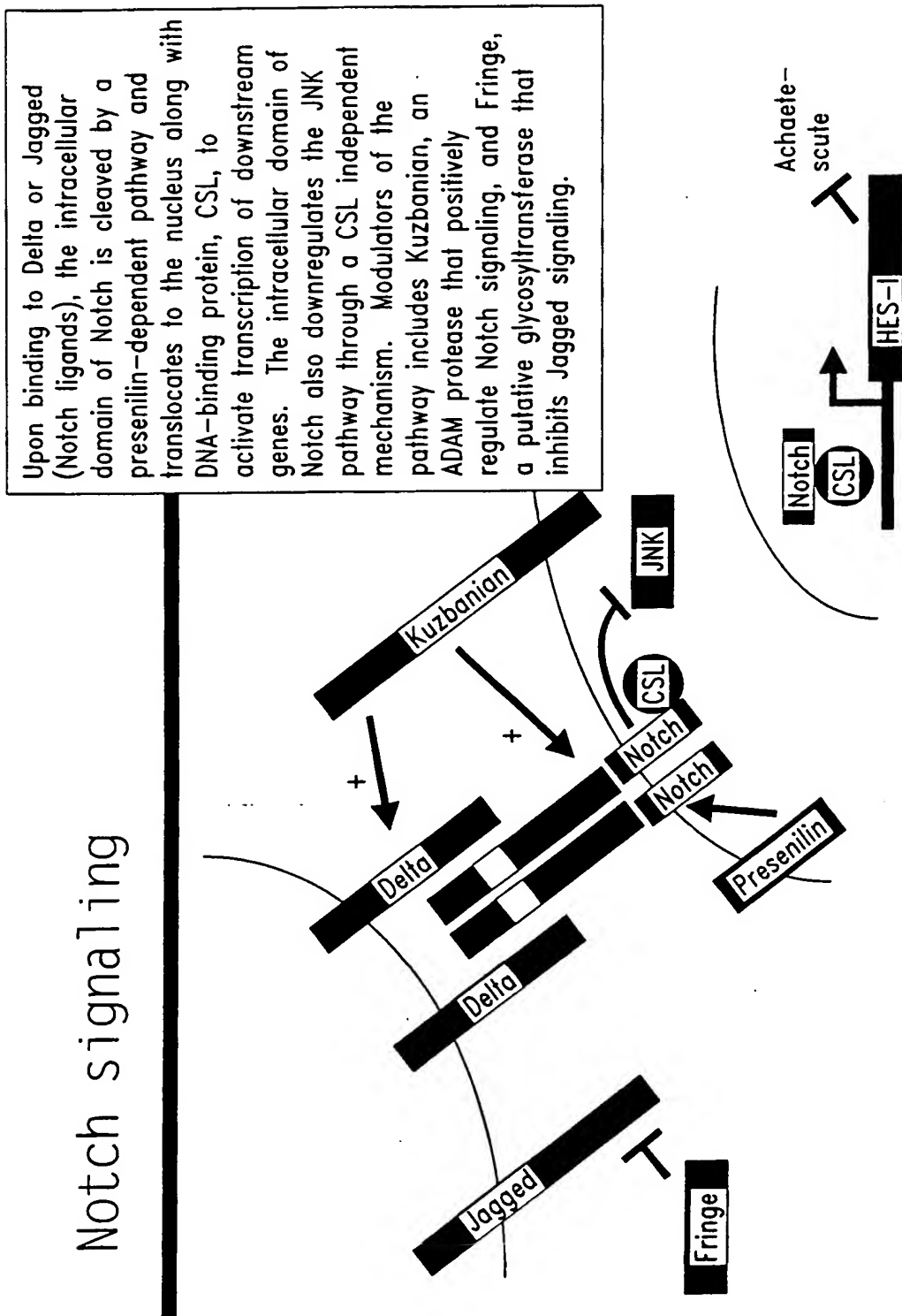


Fig. 7